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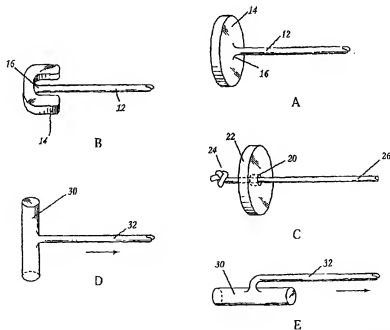
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(54) Title: DEVICES AND METHODS TO TREAT A PATIENT



(57) Abstract: Disclosed are methods and apparatus for implantation into the walls or an organ such as the stomach. Deformable or inflatable anchors with a connector between are used to pull the walls of the organ together, or to implant devices in the wall of the organ. Also disclosed are surgical instruments useful in practicing the disclosed methods.



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DEVICES AND METHODS TO TREAT A PATIENT

Related Applications

The present application claims priority to U.S. Provisional Patent Application Serial No. 60/556,004 filed March 23, 2004 by Michael Gertner, M.D., entitled "BARIATRIC DEVICES AND IMPLANTATION METHODS," to U.S. Provisional Patent Application Serial No. 60/584,219 filed July 1, 2004 by Michael Gertner, M.D., entitled "DEVICES AND METHODS FOR PERCUTANEOUS GASTROPLASTY," to U.S. Provisional Patent Application Serial No. 60/603,944 filed August 23, 2004 by Michael Gertner, M.D., entitled "DEVICES AND METHODS TO TREAT MORBID OBESITY," and to U.S. non-Provisional Application Serial No. 10/974,248 filed October 27, 2004, entitled "DEVICES AND METHODS TO TREAT A PATIENT," all of which are herein incorporated by reference in their entirety.

15 Background of the Invention

Field of the Invention

The present invention relates to methods and apparatus for implanting devices in the wall of a hollow organ, including devices to reduce the volume of the stomach.

Description of the Related Art

20 Obesity is a public health problem of growing national and international importance. There are an estimated 60 million obese adults and 2 million obese adolescents in the United States as of 2004. By some estimates, there are 1 billion obese individuals worldwide.

Over the past four decades, there have been numerous surgical procedures and
25 devices developed to treat the morbidly obese. In general, there are two physiologic
aspects of all past and current procedures: malabsorption and mechanical
restriction/volume reduction.

Many of the procedures performed in the past have proven to be impractical, dangerous, or detrimental to the health of the patients and are now of historical importance only. An example of a failed procedure is the jejunal-ileo bypass in which a malabsorptive

state is created through the bypass of a large portion of the intestine through the creation of a surgical anastomosis between the jejunum and the ileum. While patients initially lost a great deal of weight, liver failure or liver damage occurred in over one-third of the patients which necessitated reversal of the surgical procedure.

5 The Roux-en-Y (The Roux) bypass operation has evolved to become the most commonly performed surgical procedure to treat the morbidly obese. It combines a small degree of malabsorption with a 90% reduction in the volume of the stomach. In the United States, 150,000 procedures are predicted for the year 2004. This number is expected to rise to 500,000 procedures by 2006. The procedure has been performed since the late
10 1970's and the long-term data has been very good. The advent of laparoscopic surgery and hence the laparoscopic Roux-en-Y bypass in combination with excellent follow-up results from the open procedure are reasons for the proliferation of the Roux procedure.

 Despite the efficacy of the Roux procedure and the recent laparoscopic improvements, it remains a highly invasive procedure with substantial morbidity including
15 a 1-2% surgical mortality, a 20-30% incidence of pulmonary morbidity such as pneumonia, pulmonary embolism, etc., and a 1-4% chance of leak at the anastomotic site which can result in a spectrum of consequences which can range from an extended hospital stay to death.

 The Roux procedure requires general anesthesia and muscle paralysis which, in the
20 morbidly obese population, is not of small consequence. There is also a substantial rate of anastomotic stricture which results in severe lifestyle changes for patients. For example, many patients are forced to vomit after meals. Furthermore, although minor when compared to previous malabsorptive procedures, the malabsorption created by the Roux-en-Y can dramatically affect the quality of life of patients who undergo the procedure.

25 The largest problem from the perspective of the patient is that the Roux-en-Y is not reversible, which dramatically limits the number of patients willing to undergo the procedure. In particular, it severely limits the number of procedures which can or should be performed on adolescents.

 Recently, minimally invasive procedures and devices which create a feeling of early
30 satiety have been introduced into the marketplace in an attempt to address some of the issues above. The LAP-BAND™ is a band which encircles the stomach at the region of

the fundus-cardia junction. It requires general anesthesia, a pneumoperitoneum, muscle paralysis, and extensive dissection of the stomach at the level the gastroesophageal junction. Although less invasive than the Roux procedure and potentially reversible, the LAP-BAND™ is nonetheless quite invasive. It also does not reduce the volume of the stomach and patients report a feeling of hunger much of the time. The banding procedure is also fraught with complications. In a recently published article (Camerini et.al. Thirteen Years of Follow-up in Patients with Adjustable Silicone Gastric Banding for Obesity: Weight Loss and Constant Rate of Late Specific Complications. Obesity Surgery, 14, 1343-1348), the authors reported 60% prevalence of late band removal secondary to complications.

Other procedures which have been tried in the past and which offer varying degrees of weight loss include any of a variety of "gastroplasty" procedures. These procedures were among the first operations attempted to control weight, the so-called "stomach stapling" procedures. These procedures were tried before the Roux-en-Y became popular and are considered to offer patients less weight loss than the Roux but also a less invasive and more reversible procedure.

The first gastroplasty procedure was the horizontal gastroplasty in which a horizontal row of staples was applied across the stomach from the lesser curvature to the fundus without dividing the stomach between the staples. After application of the staple line, several of the staples are removed (in the middle of staple line) leaving an os, or opening, for flow of food. Although this procedure resulted in weight loss, in most patients, the staples pulled through the walls of the stomach leading to an unwanted reversal of the plasty procedure. Furthermore, even before the staples pulled through the walls of the stomach, the os would dilate, allowing for progressively larger amounts of food to be swallowed by the patient.

Following, and overlapping with the horizontal gastroplasty, a procedure was developed called the vertical banded gastroplasty, or VBG, which again, created a restricting os for food. In the VBG, the border of the os is the lesser curvature of the stomach which is less apt to dilate than the center portion of the stomach. Furthermore, the procedure completely excludes the fundus; the fundus is thought to also easily dilate and is actually physiologically "programmed" to dilate during feeding..."receptive

relaxation." A recent prospective, randomized trial compared the VBG to the adjustable banding procedure (see below) and found a highly significant advantage of the VBG over the adjustable banding procedure (reference). Although in this study the VBG was performed laparoscopically, the laparoscopic VBG procedure is difficult to perform because the procedure is not standardized and a "tool box" does not exist for the surgeon to carry out the procedure; furthermore, the procedure is not a reversible one.

The Magenstrasse and Mill (M&M) procedure is an evolving gastroplasty technique wherein the greater curvature of the stomach is essentially taken out of the path of food, leaving a tube of stomach, the Magenstrasse, or "street of the stomach," which is comprised of the lesser curvature. This procedure is similar to the VBF except that the longitudinal staple line of the stomach extends further along the lesser curvature and into the antrum. Furthermore, a PTFE wrap is not required at the point where an EEA stapler is placed through the stomach. The antrum is preserved in this procedure. The theory behind leaving the antral "mill" is that it will continue to serve its normal function of mixing, grinding, retropulsion, and well-regulated expulsion of chyme into the duodenum. An authoritative study on the operation is incorporated herein by reference (Johnston et. al. The Magenstrasse and Mill Operation for Morbid Obesity; Obesity Surgery 13, 10-16).

A recent Meta-analysis and systematic review (Buchwald et. al. Bariatric Surgery: A Systematic Review and Meta-analysis. JAMA vol. 292, no 14. pps 1724-1737) indicated that vertical gastroplasty (avg. excess weight loss of 68.2%) is superior to adjustable banding (avg excess weight loss of 47.5%) and gastric bypass (avg excess weight loss of 61.6%). In another study, this one a single prospective and randomized study, laparoscopic vertical banded gastroplasty was compared directly with laparoscopic banding (Morino et. al. Laparoscopic Adjustable Silicone Gastric Banding Versus Vertical Banded Gastroplasty in Morbidly Obese Patients. Annals of Surgery. Vol. 238 (6) pps. 835-842). In the gastroplasty group, excess weight loss after three years was 58.9% and 39% in the banding group. In the gastroplasty group, late reoperation was 0% and 24.5% in the banding group.

In summary, the vertical gastroplasty procedure appears to be superior to the banding procedure. However, the gastroplasty procedure is not easily performed

laparoscopically; furthermore, it is not reversible because the stomach is divided. If the stomach is not divided, then the staples will eventually "pull through" the wall of the stomach and the patients will again gain weight again. Division of the stomach was one of the early lessons which was determined to be necessary for the gastroplasty procedure to successful long-term. Therefore, there is a need to standardize the verticle banded gastroplasty procedure and create a procedure which is both easy to perform and reversible.

The intragastric balloon is not a new concept. The intragastric balloon is meant to displace volume within the stomach such that a smaller volume of food leads to an earlier feeling of satiety. Currently, intragastric balloons on the market are not fixed to the stomach. As a consequence, the intragastric balloons lead to complications such as obstruction and mucosal erosion. As a consequence, the balloons are removed after a maximum of six months. A balloon which is fixed to the wall of the stomach could potentially improve the intragastric balloon device and allow longer term implantation. In a prospective, non-randomized, unblinded study (Sallet et. al. Brazilian Multicenter Study of the Intragastric Balloon; Obesity Surgery, 14, 991-998), the average excess weight loss was 48.3% after 1 year. However, the incidence of nausea and vomiting was 40% and epigastric pain was 20%; balloon impaction occurred in .6% of patients.

Percutaneous Endoscopic Gastrostomy (PEG) refers to a procedure in which a gastrocutaneous tract is created using percutaneous means. A recent update of the procedure can be found on the Society of American Gastrointestinal Endoscopic Surgeons (SAGES) website, and is incorporated herein by reference. Briefly, the procedure involves insufflation of the stomach with and under visualization with an endoscope. A small incision is made in the skin and a needle is advanced into the stomach (the stomach sits just under the abdominal wall when insufflated) under endoscopic visualization. A feeding tube is then placed over the needle to create a gastrocutaneous tract with the feeding tube inside the tract. The feeding tube is secured with an external bolster. Over the ensuing weeks, a permanent tract evolves between the stomach mucosa and epithelium of the skin, after which, the bolster can be removed without consequence. When the feeding tube is to be removed, the gastrocutaneous tract will close on its own as food will preferentially be

delivered antegrade (the path of least resistance) to the duodenum, thereby allowing the tract to heal.

More recently, there has been an effort to develop even less invasive devices and procedures which do not involve incisions at all. For the most part, these procedures are performed from within the stomach with an endoscope and by a physician with a high degree of endoscopic skill. For example, U.S. Patent No. 6,558,400 describes methods and devices to create partitions in the stomach. Fasteners or staplers applied through an endoscope from within the stomach are used to accomplish the partitions. Similarly, U.S. Patent Application Publication No. 2004/0122456 describes another set of methods and devices to reduce the volume of the stomach. Expandable anchors are deployed both on the anterior and posterior wall of the stomach by way of an endoscope. Flexible sutures are brought out of the patient's mouth and the sutures are crimped together within the stomach in order to bring the walls of the stomach closer together. Patent application WO2004/004542 describes a device which is advanced through an endoscope and grasps or applies suction to a fold of mucosa to apply fasteners through the mucosal and serosal layers of the stomach.

Endoscopic procedures to manipulate the stomach are time consuming because of the technical difficulty of the endoscopy; they also require a large endoscope through which many instruments need to be placed for these complex procedures. Due to the large girth of the endoscope, patients typically will require general anesthesia, limiting the "non-invasive" aspects of the procedure. Furthermore, the procedures require advanced endoscopic skill which would need to be acquired by most practitioners. Such skill adaptation can take a significant amount of time, which will limit adoption of the procedure by the physician community. A further issue is that there is a limitation on the size of the anchors which can be placed because the endoscope has a maximum size.

Summary of the Invention

One embodiment of the present invention is a method for reducing the interior volume of an organ comprising passing a first end of a first surgical instrument through the patient's skin, through a first exterior surface of the organ, through the interior of the organ, and then through a second exterior surface of the organ, so that the surgical instrument

traverses the organ, deploying a first anchor from the surgical instrument wherein the first anchor is located adjacent to the second exterior surface of the organ, partially or completely withdrawing the surgical instrument, deploying a second anchor wherein the second anchor is located adjacent to the first exterior surface of the organ, providing a
5 connector between the first and second anchors, wherein the length of the connector between the first and second anchors is such that the first and second anchors urge the first and second exterior surfaces of the organ toward each other, thereby reducing the volume of the organ. In some embodiments, the first and second anchors are deployed from the same surgical instrument. Another aspect of the invention is a method of reversing the volume
10 reducing procedure by cutting or otherwise dividing the one or more connectors between the first and second anchors. In another embodiment of the method, the organ is a gastrointestinal organ. The method may further comprise creating a space outside the organ adjacent to the second exterior surface thereof by introducing a volume-filling substance into a potential space adjoining the second exterior surface. The potential space can be
15 expanded by the injection of a gas, liquid, gel, foam, or solid, into the potential space, by the inflation of a balloon placed in the potential space, or by blunt dissection. The method may further comprise insufflating the organ prior to the passing of the first end of a surgical instrument through a first exterior surface of the organ. In some embodiments, the patient's skin is overlying the patient's stomach, the organ is the patient's stomach, the
20 first exterior surface is the anterior wall of the stomach, the second exterior surface is the posterior wall of the stomach, and the potential space is the lesser sac of the peritoneum. The method may further comprise urging the anterior and posterior walls of the stomach closer together by shortening the length of the connector between the first and the second anchors. In some embodiments, the surgical instrument is inserted into the patient's
25 abdomen by directly penetrating the patient's skin and abdominal wall, by passing the surgical instrument through a laparoscopic port, or by passing the surgical instrument through an incision in the patient's skin and abdominal wall.

Another embodiment of the invention is a method for reducing the volume of a patient's stomach, comprising passing a first anchor through a patient's abdominal skin,
30 and through the patient's anterior and posterior stomach walls, wherein the first anchor is in a reduced profile configuration, passing a second anchor through a patient's abdominal

skin and adjacent to the patient's anterior stomach wall, connecting the first and second anchors by means of a connector that passes through the stomach, urging the first and second anchors toward each other, and holding the anterior and posterior walls of the stomach together with the first and second anchors, wherein the first anchor is in a
5 deployed configuration and the connector prevents the first and second anchors from moving apart. Some embodiments further comprise passing the second anchor through the patient's abdominal skin while the second anchor is in a reduced profile configuration, as well as the second anchor subsequently being in a deployed configuration. In some
10 embodiments, the holding of the anterior and posterior walls of the stomach together is performed in a manner that permits some space between mucosal surfaces of the stomach interior of the walls. In other embodiments it is performed in a manner that permits contact between mucosal surfaces of the stomach interior of the walls. In some
embodiments, the holding step is performed by adjusting the relative position of the connector and the second anchor, and then configuring the second anchor to engage the
15 connector in a manner that prevents the first and second anchors from moving apart.

Another embodiment of the invention is a fastening assembly, comprising a first anchor, a second anchor, and a connector, wherein the first anchor comprises a relatively planar body attached to the connector, the body of the first anchor having a relatively
20 planar deployed profile and a reduced profile configuration, wherein the second anchor comprises a relatively planar body, a hole or other passageway approximately in the center of the body of sufficient diameter to allow passage of the connector through the hole or other passageway, one, two or more gripping elements projecting into the hole or other
passageway, and one, two or more attachment structures accessible from a top surface of the body, the body of the second anchor having a relatively planar deployed profile and a
25 reduced profile configuration, and wherein the gripping elements prevent movement of the second anchor along the longitudinal axis of the connector in the direction away from the first anchor when the connector is disposed in the hole or other passageway when the second anchor is in its deployed configuration.

Another embodiment of the invention is a biocompatible surgical anchor adapted for
30 use in connecting the walls of a hollow organ or anchoring a device to the wall of a hollow organ, comprising a body portion, wherein the body portion is compressible and can

assume a relatively planar deployed configuration and a reduced-profile compressed configuration, wherein the body portion has an opening extending therethrough, and a gripper on the body portion adapted to grip an elongated connector as such a connector extends through the opening, wherein the gripper engages such a connector when the anchor is in the deployed configuration, and releases such a connector when the anchor is in the compressed configuration. In some embodiments, the body portion is inflatable and can assume a relatively planar deployed configuration when inflated and a reduced-profile configuration when uninflated, wherein the body portion has an opening extending therethrough, and a gripper on the body portion adapted to grip an elongated connector as such a connector extends through the opening, wherein the gripper engages such a connector when the anchor is inflated, and releases such a connector when the anchor is uninflated.

Another embodiment of the invention is a fastening assembly, comprising a first anchor, a second anchor, and a connector, wherein the first anchor comprises an inflatable body attached to the connector, the body of the first anchor having a deployed configuration when inflated that is relatively spherical and a reduced profile configuration when uninflated wherein it is readily deformable, wherein the connector is hollow and configured to allow inflation of the first anchor by delivery of a filling substance through the connector to the first anchor.

Another embodiment of the invention is a method of fastening a device to one wall of a gastrointestinal organ comprising passing said device through a patient's abdominal skin, and through the patient's anterior stomach wall, wherein the device is in an undeployed configuration, passing an anchor through a patient's abdominal skin wherein the anchor is in an undeployed configuration, connecting the device and the anchor by means of a connector that passes through the wall of the gastrointestinal organ wherein the connector prevents the device and second anchor from moving apart, and deploying the anchor to its deployed configuration. In another embodiment, the device comprises an inflatable body attached to the connector, the body of the device having a deployed configuration when inflated that is relatively spherical and a reduced profile configuration when uninflated wherein it is readily deformable, wherein the connector is hollow and

configured to allow inflation of the device by delivery of a filling substance through the connector to the device.

In another embodiment, the device is an electrical stimulator.

In yet another embodiment, the device is adapted to deliver a medicine.

5 In any of the embodiments, the reduced profile configuration can be substantially compressed and/or uninflated in the undeployed configuration, or inflated and/or uncompressed, in the deployed configuration.

10 In any of the embodiments, it can be the case that the gripping elements do not prevent movement of the second anchor along the longitudinal axis in either direction when the connector is disposed in the hole or other passageway when the second anchor is in its reduced profile configuration.

15 In any of the embodiments it may be the case that the gripper or gripping elements, when engaging a connector, allow movement of the connector in one direction relative to the anchor but prevents movement of the connector in an opposite direction relative to the anchor.

20 In any of the embodiments, the body of the first anchor can be inflatable, have a relatively planar deployed profile when inflated and a reduced profile configuration when uninflated, be readily deformable when uninflated, and the connector can be hollow and configured to allow inflation of the first anchor by delivery of a filling substance through the connector to the first anchor.

25 In any of the embodiments, the body of the second anchor can be inflatable, wherein the second anchor has a relatively planar deployed profile when inflated and a reduced profile configuration when uninflated, and wherein the second anchor is readily deformable when uninflated.

30 In any of the embodiments it may be the case that the anchor has an inflation tube connected to the body portion configured to allow inflation of the body portion by delivery of a filling substance through the inflation tube to the body portion.

In any of the embodiments, it can be the case that the filling substance hardens, cures, polymerizes, or become a gel over time, and is optionally bioabsorbable with further time.

In any of the embodiments, it can be the case that the second anchor has one or more attachment structures accessible from a top surface of the body.

In any of the embodiments, the first anchor, the second anchor, or both the first and the second anchors, can deliver an electrical signal to tissue when placed in contact with the tissue. In some embodiments, only the first or second anchor is placed which then delivers an electrical signal.

Another embodiment of the invention is a surgical instrument comprising a proximal end, a distal end, and a shaft with a channel, the distal end comprising a tissue penetrator adapted to penetrate, cut, or dilate the wall of a hollow organ, and a tissue grasper adapted to grasp a wall of a hollow organ, an expander within the channel of the surgical instrument wherein the expander can be deployed from within the channel of the instrument and extend through a wall of the hollow organ while the wall is being grasped by the tissue grasper, and is adapted to introduce a volume-filling material or device beyond the hollow organ.

Another embodiment of the invention is a surgical instrument comprising a proximal end, a distal end, and a channel, wherein the distal end comprises a tip which has an open configuration and a closed configuration, wherein in the closed configuration, the tip is adapted to penetrate a wall of a hollow organ, wherein in the open configuration, the tip can grasp the wall of the hollow organ, and wherein the channel comprises a diameter of about 2 mm to 12 mm.

Another embodiment of the invention is an anchor implantation instrument comprising, an outer sheath, a middle sleeve disposed within the outer sheath, an inner member disposed within the middle sleeve, and a channel within the inner member, wherein the middle sleeve has an outer diameter and a longitudinal axis, and the outer diameter of the middle sleeve is such that it can slide within the outer sheath along the longitudinal axis of the middle sleeve, wherein the inner member has an outer diameter, and the outer diameter of the inner member is such that it can slide within the middle sleeve along the longitudinal axis of the middle sleeve, wherein the inner member has a grasper suitable for grasping a foldable anchor and wherein the outer sheath has a distal portion which has an inner diameter sufficient to hold a foldable anchor in its folded configuration.

One aspect of the invention is a surgical device system that includes a laparoscopic instrument and a grasping mechanism. The laparoscopic instrument includes a fastener adapted to receive a connector having a proximal end and a distal end. In one embodiment, the grasping mechanism is positioned on the instrument. Alternatively, the grasping mechanism is positioned on the connector.

The grasping mechanism, in one embodiment of the invention, includes an actuator that is controllable at the proximal end of the connector or the instrument. The actuator may be adapted to activate the grasping mechanism to reversibly engage the fastener. In one embodiment, the connector includes a removable outer sheath.

Another aspect of the invention includes a method of treating a patient, including positioning a first surgical device that has a reversibly engaged fastener near or against the patient's organ, positioning or forming an elongate member in the patient, where the elongate member includes a proximal end, a distal end and a circumferential wall forming a lumen sized to receive a connector therethrough, and where the connector has a proximal end and a distal end, engaging the first fastener with the connector, then advancing the connector through the lumen of the elongate member, the elongate member thence percutaneously applied through a first portion of the organ and then through a second portion of the organ, releasing the first fastener from the first surgical device such that the first fastener remains engaged with the connector, advancing a second surgical device through the lumen, wherein the second surgical device includes a second fastener configured to track the connector, urging the first and second portions of the organ toward each other with the first and second fasteners, releasing the second fastener from the second surgical instrument such that the second fastener engages the connector, and cutting the connector such that the connector has a length that spans the first and second fasteners.

In one embodiment the organ is the patient's stomach. In another embodiment, urging the first and second portions of the organ reduces the volume of the organ and creates a resistance to the flow of material into the organ. In a related embodiment, the urging causes mucosa of the first and second portions of the patient's stomach to contact each other.

In yet another embodiment, the elongate member is a laparoscopic port that is positioned in the patient. The elongate member and the connector, in one embodiment of the invention, are one device.

5

Brief Description of the Drawings

Figures 1A-1E are perspective views of embodiments of the posterior anchor and connector.

Figures 1F and 1G are side views of an inflatable embodiment of posterior anchor and connector.

10 Figures 2A and 2B are a perspective view and top view of one embodiment of an anterior anchor, respectively.

Figures 2C and 2D are side sectional views of the embodiment of the anterior anchor of FIGs. 2A and 2B, taken along the line B-B in FIG. 2B, in its deployed and reduced profile configuration, respectively.

15 Figures 2E and 2F are side sectional views of another embodiment of an anterior anchor, taken along the same line as FIGs. 2C and 2D, in its deployed and reduced profile configuration, respectively.

Figures 2G is a perspective view of an inflatable embodiment of an anterior anchor.

20 Figures 2H and 2I are side sectional views of the embodiment of the anterior anchor of FIG. 2G, taken along the line D-D in FIG. 2G, in its deployed and reduced profile configuration, respectively.

Figure 3A is a perspective view of another embodiment of an anterior anchor.

Figures 3B and 3C are perspective views of the embodiment of the anterior anchor shown in FIG. 3A in its reduced profile and deployed configuration, respectively.

25 Figure 3D is a perspective view of another embodiment of an anterior anchor.

Figures 4A and 4A' are a side and blow-up view, respectively, of one embodiment of a tissue grasping instrument with the distal end in its open configuration.

Figures 4B and 4B' are a perspective and blow-up view, respectively, of the tissue grasping instrument of FIG. 4A with the distal end in its closed configuration.

Figures 4C and 4C' are a perspective and blow-up view, respectively, of another embodiment of the tissue grasping instrument with the distal end in its closed configuration.

Figures 5A is a side view of one embodiment of an anchor implantation instrument.

5 Figure 5B is a perspective view of the distal end of the anchor implantation instrument of FIG. 5A and an anterior anchor and connector.

Figure 5C is a side sectional view of the distal end of the anchor implantation instrument of FIGs. 5A and 5B, taken along line C-C in FIG. 5B, with the anterior anchor in its reduced profile configuration.

10 Figure 6A illustrates the first step in one embodiment of a method of reducing the volume of the stomach. Shown is a side sectional view of a patient's abdomen with the instrument of FIG. 4 inserted into the patient's abdomen through a laparoscopic port.

Figure 6B illustrates the next step in one embodiment of a method of reducing the volume of the stomach. Shown is a side sectional view of a patient's abdomen with the instrument of FIG. 4 grasping the posterior wall of the stomach and a needle being inserted
15 into the potential space of the lesser peritoneal sac.

Figure 6C illustrates the next step in one embodiment of a method of reducing the volume of the stomach. Shown is a side sectional view of a patient's abdomen with the instrument of FIG. 4 grasping the posterior wall of the stomach and a posterior anchor and
20 connector deployed in the expanded potential space of the lesser peritoneal sac.

Figure 6D illustrates the next step in one embodiment of a method of reducing the volume of the stomach. Shown is a side sectional view of a patient's abdomen with a posterior anchor and connector deployed in the expanded potential space of the lesser peritoneal sac, with the connector passing out of the patient's abdomen through a
25 laparoscopic port.

Figure 7A illustrates the next step in one embodiment of a method of reducing the volume of the stomach. Shown is a side sectional view of a patient's abdomen with the instrument of FIG. 5C placing an anterior anchor in the patient's abdomen adjacent to the anterior wall of the stomach.

30 Figure 7B illustrates the next step in one embodiment of a method of reducing the volume of the stomach. Shown is a side sectional view of a patient's abdomen with an

anterior anchor in its deployed configuration on the connector, with the anterior and posterior walls of the stomach urged together.

Figure 7C illustrates the next step in one embodiment of a method of reducing the volume of the stomach. Shown is a side sectional view of a patient's abdomen after the
5 connector has been cut flush with the anterior anchor.

Figure 8A illustrates an embodiment of a method of reducing the volume of the stomach. Shown is a side sectional view of a patient's abdomen after two posterior anchors and connectors have been deployed adjacent to the posterior wall of the stomach, with the connectors passing out of the patient's abdomen through laparoscopic ports.

10 Figure 8B shows the connectors of FIG. 8A with clamps placed on the connectors outside the patient's body to temporarily hold the connectors in a test position.

Figure 9 is a perspective view showing three transgastric fastening assemblies deployed in a patient's stomach.

Figure 10A illustrates one embodiment of a method for deploying a volume
15 displacing device in the stomach. Shown is a side sectional view of a patient's abdomen after an uninflated balloon anchor has been inserted inside the patient's stomach with a connector passing out of the stomach, through the anterior stomach wall, and a laparoscopic port.

Figure 10B illustrates one embodiment of a method for deploying a volume
20 displacing device in the stomach. Shown is a side sectional view of a patient's abdomen with the balloon anchor in its deployed position, held in place by an anterior anchor.

Figure 11A illustrates a volume displacing device which resides outside the stomach and is shown in an undeployed state.

Figure 11B depicts a volume displacing device which resides outside that stomach
25 and is shown in a deployed state.

Figure 11C depicts a volume displacing device which resides outside the stomach and is fixed to the anterior wall of the stomach.

Figure 12 depicts a fastener which is applied in a laparoscopic procedure. Shown is
30 a connector being pushed through the walls of the stomach to meet the fastener component which is applied through the laparoscopic ports and which meets the connector portion.

Figure 13 depicts the second step in the laparoscopic procedure in which the anterior fastener is brought toward the posterior fastener over the connector.

Figure 14 depicts the third step in the laparoscopic procedure in which the anterior and posterior walls of the stomach are urged together.

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Detailed Description of the Preferred Embodiment

Anatomy of the Stomach

The region behind the stomach is referred to as the lesser peritoneal sac. It is a potential space between the retroperitoneum and the posterior wall of the stomach. To the left of the midline, the posterior wall of the stomach is generally free from the peritoneal surface of the retroperitoneum. To the right of the midline, the posterior wall of the stomach is more adherent to the retroperitoneum although the adherence is generally loose and the adhesions can be broken up rather easily with gentle dissection.

The stomach is comprised of several layers. The inner layer is the mucosa. The next layer is the submucosa followed by the outer muscular layers. Surrounding the muscular layers is the serosal layer. This layer is important with regard to implants and healing because it is the adhesive layer of the stomach; that is, it is the layer which, when breached, heals with scar tissue formation. Implants adhering to this layer are less likely to migrate into the stomach. Reference to "stomach wall" or "wall of the stomach" as used herein include the entire thickness of the stomach, including the mucosa, submucosa, muscular layers, and serosa. The "anterior wall of the stomach" is the portion of the stomach closest to the muscular abdominal wall and the "posterior wall of the stomach" is the part of the stomach closest to the retroperitoneum.

"Transgastric fastening assembly" refers to a permanent or semi-permanent implant and comprises at least one posterior anchor, at least one anterior anchor, and a fastener to connect the posterior and anterior anchors. The "fastener" can refer to any means of connection including but not limited to a material connection, an electromagnetic connection, or a chemical connection. As used herein, a "connector" is a fastener used to materially connect anterior and posterior anchors. As used herein, the "posterior anchor" is the anchor in a preferred embodiment which is adjacent to the posterior wall of the

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stomach when deployed. The “anterior anchor” is the anchor in a preferred embodiment which is approximated to the anterior wall of the stomach when deployed.

As used herein when referring to portions of a surgical instrument, “proximal” refers to the end of the instrument which is closest to the surgeon when the instrument is used for its intended purpose, and “distal” refers to the end of the instrument which is closest to the patient when the instrument is used for its intended purpose. When used to refer to the gastrointestinal tract, “proximal” is toward the mouth and “distal” is toward the anus.

Percutaneous surgery typically means that the procedure is performed under visualization (e.g. fluoroscopic, MRI, CAT Scan, Ultrasound, Endoscopic) which is not direct visualization and which requires an incision, typically small, somewhere in the skin. More recently, the boundaries of percutaneous surgery have been blurred as some procedures involve both direct visualization, such as with a laparoscope, and percutaneous methodology. An example of such a procedure is the laparoscopic placement of a jejunostomy or gastric feeding tube. Laparoscopic methods require general anesthesia to paralyze the abdominal muscles so that the abdomen can be filled with gas. Consequently, general anesthesia is required so as to enable paralysis of the diaphragm and respiratory muscles. A purely percutaneous procedure does not typically require general anesthesia.

As used herein, “percutaneous procedure” refers to a procedure wherein general anesthesia and general pneumoperitoneum are not used or the procedure utilizes incisions through the skin of the abdomen for access to the surgical site and not for visualization. Therefore, as used herein, percutaneous surgery and laparoscopic surgery are mutually exclusive. In the preferred embodiment, the methods described herein are performed percutaneously, although laparoscopic methods are contemplated in other embodiments; in still other embodiments, combination procedures are contemplated and in these embodiments, laparoscopic surgery is used to represent the combination of methods and devices.

STRUCTURES

Transgastric Fastening Assembly

Referring to FIGs. 1A and 1B, one embodiment of the posterior anchor 14 and connector 12 are shown in a deployed configuration (FIG. 1A), and reduced profile configuration (FIG. 1B). The connector 12 is preferably made of a biocompatible semi-rigid polymer, but it can be made from various kinds of suitable biocompatible materials known to those of skill in the art including metals, such as titanium and platinum, metal alloys, such as stainless steel, nickel-titanium, and cobalt-chromium, man-made polymers, such as polyurethane, silicone elastomers, polyglycolic acid, polylactic acid, poly (-caprolactone), polyvinylidene fluoride (PVDF), PTFE, FEP, polypropylene, or natural fibers such as silk. These materials can be used singly or in combination. For example, one portion of the connector may be bioabsorbable and another portion of the connector may be permanent. The connector 12 can vary in thickness, shape, and rigidity. For example, in the embodiment shown in FIG. 1A, the connector 12 is substantially rod-shaped, with a circular cross-section, and is semi-rigid. Those of skill in the art will recognize that the cross-section of the connector can be any of a number of shapes, such as square, hexagonal, oval, etc. In other embodiments, the connector 12 is thin and flexible, such as a surgical suture, and in still others it is rigid.

In a preferred embodiment, the posterior anchor 14 is made from a biocompatible, radio-opaque or magneto-opaque semi-rigid polymer; it can also be made from various kinds of suitable materials known to those of skill in the art including metals, metal alloys, plastics, natural materials or combinations thereof as discussed above. The posterior anchor 14 can be solid, or alternatively, can be porous, mesh-like, lattice-like, or umbrella-like. In a preferred embodiment, the posterior anchor is porous or has a porous mesh attached to it to encourage fibrous ingrowth such that it becomes permanently attached to the stomach or intestinal wall. Coatings can be added to the anchor to encourage tissue ingrowth. In other embodiments, the posterior anchor is solid and treated to discourage tissue ingrowth. In other embodiments, the anterior anchor has a xenograft or allograft material attached to the anchor. In a preferred embodiment, the posterior anchor 14 is disc-shaped, but those of skill in the art will recognize that other embodiments are possible, such as those shown in FIGs. 1C and 1D, or disclosed in U.S. Patent Application Publication No. 2004/0122456 which is herein incorporated by reference; note particularly

the description of anchor structures. The posterior anchor can also be bioabsorbable in whole or in part in some embodiments.

In the embodiment shown in FIGs. 1A and 1B, the connector 12 is fastened to the posterior anchor 14 at an attachment point 16 which is preferably a permanent, e.g. welded or molded, connection. Such a weld or connection can comprise, for example, a thermoformed polymer, a metallic weld, or a molded or other integral structure. In a preferred embodiment, a biocompatible thermoformed polymer is used because of its flexibility and ability to yield to the continuous motion of the stomach. More preferably, the connector and posterior anchor are produced as a single, continuous injection molded component.

Other suitable means of fastening the connector to the posterior anchor are also contemplated and do not necessarily result in a connector and posterior anchor becoming permanently attached. For example, in one embodiment shown in FIG. 1C, one end of the connector is passed through a hole 20 near the center of the posterior anchor 22, and a stop 24, such as a knot or enlarged molded region, is formed on the end of the connector to prevent its passage back through the hole in the posterior anchor. In this embodiment, the posterior anchor 22 can be free to move along the length of the connector 26, but is prevented from being removed from one end of the connector by the stop 24.

In the embodiment shown in FIGs. 1A and 1B, the posterior anchor 14 preferably has a deployed configuration (FIG. 1A), and reduced profile configuration (FIG. 1B). The posterior anchor 14 can be deformed to a folded configuration wherein its profile is reduced to facilitate insertion of the anchor through the walls of the stomach or other tissue as described in more detail below. In one embodiment, the posterior anchor 14 is made of a semi-flexible material having shape memory, so that once the anchor is deployed within the patient, it will return to its original shape shown in FIG. 1A, preventing it from being easily pulled back through the tissue. Preferably, the posterior anchor is inflatable in place of, or in addition to, having shape memory, which allows for a much larger deployed profile relative to its undeployed profile. (See below).

Figures 1D and 1E show an alternative embodiment of the posterior anchor 30 and connector 32 in a deployed configuration (FIG. 1D), and a reduced profile configuration (FIG. 1E). In this embodiment, the posterior anchor 30 is elongated, having major and

minor dimensions, and preferably having a rod or bar shape. By aligning the connector 32 substantially parallel to the posterior anchor 30, its profile is reduced to facilitate insertion of the anchor through the walls of the stomach or other tissue. When the anchor leaves its surrounding sheath (see below), tension on the connector 32 in the direction of the arrow in FIG. 1E will urge the posterior anchor 30 into a substantially perpendicular orientation relative to the connector 32, as shown in FIG. 1D, preventing it from easily being pulled back through the tissue. The connection between the posterior anchor 30 and the connector 32 can be hinged. Alternatively, the connector 32 can be made of a semi-rigid material which is permanently connected or welded to the posterior anchor 30. If the connector is deformed to a bent position, shown in FIG. 1E, it will return to its original straight shape shown in FIG. 1D once the anchor is deployed within the patient, preventing the posterior anchor from easily being pulled back through the tissue. This anchor 30 can be inflatable as well, which allows for a much larger deployed profile relative to its undeployed profile.

In a preferred embodiment, shown in FIGs. 1F and 1G, the posterior anchor is inflatable. The anchor has an inflatable disc-shaped body 34 which is readily deformable when in its reduced profile (i.e., uninflated) configuration as shown in FIG. 1F. In the preferred embodiment, the posterior anchor body 34 is disc-shaped, but those of skill in the art will recognize that other embodiments are possible, such as those shown in FIGs. 1C and 1D, or in which the inflatable anchors are square shaped, rectangular, or amorphous, or have a shape disclosed in U.S. Patent Application Publication No. 2004/0122456 which is herein incorporated by reference; note particularly the description of anchor structures. The body can be inflated with a substance delivered through a hollow connector 35. When the interior space 36 of the anchor body is inflated, the anchor assumes its deployed configuration shown in FIG. 1G.

The inflatable posterior anchor can have a valve 38 located between the anchor body 34 and the connector 35. Alternatively, the valve is located in the portion of the connector located outside the patient, the valve (e.g. stopcock type valve) being controlled by the operator until the anterior anchor is placed (see below). In this alternative embodiment, the filling substance is trapped in the posterior anchor after the anterior anchor is deployed and the connector is cut and sealed, preferably flush with the anterior

anchor (see below). The filling substance can be a gas, liquid, or material which changes phase with time (i.e. it may harden, cure, polymerize, or become a gel with time). Preferably, the surface of the posterior anchor adjacent to the posterior wall of the stomach has a mesh fixed to it to encourage tissue ingrowth. In some embodiments, part, or all of the anchor material is comprised of a biodegradable material.

Figures 2A (perspective view) and 2B (plan view) show an embodiment of the anterior anchor 40. The anterior anchor has a disc-shaped body 42 with a hole or other passageway 44 substantially in the middle of the body. Two gripping elements 46 project into the center of the hole or other passageway. With respect to the gripping elements, there can be as few as one or more than two. The gripping elements may have teeth 50 angled toward the top surface of the anchor. Optionally, two hooks 52, or other graspable recesses, appendages, or structures, are located on the top surface of the anterior anchor. Hooks 52 allow for attachment of a surgical instrument during deployment of the anterior anchor in the patient as described below. Alternatively, there can be none, one, or more than two graspable recesses, appendages, or structures on the top surface of the anchor. In the preferred embodiment, the anterior anchor body 42 is disc-shaped, but those of skill in the art will recognize that other embodiments are possible, as disclosed in U.S. Patent Application Publication No. 2004/0122456 which is herein incorporated by reference; note particularly the description of anchor structures.

Figures 2C and 2D are cross sections of the anterior anchor of FIGS. 2A and 2B, taken along the line B-B in FIG. 2B. Figure 2C shows the anterior anchor in its deployed configuration with the connector 12 of FIG. 1A passing through the hole or other passageway 44 in the body of the anchor. In the deployed configuration, the gripping elements 46 and teeth 50 engage the connector 12 with sufficient pressure to prevent movement of the anchor along the connector 12 in the direction of the arrow in FIG. 2C, which would increase the distance between the anterior anchor and posterior anchor (not shown). In FIG. 2D, the anterior anchor 40 is in its reduced profile configuration with the connector 12 of FIG. 1A passing through the hole or other passageway 44 in the body of the anchor. Preferably, the anterior anchor is made of a semi-rigid polymer which allows the anchor to be deformed into a substantially folded configuration illustrated in FIG. 2D. When in this configuration, the gripping elements 46 and teeth 50 do not significantly

engage the connector 12. This allows movement of the anterior anchor 40 along the length of the connector 12 in the directions illustrated by the arrows in FIG. 2D. Once the anterior anchor is in the desired position along the connector 12, the anterior anchor is permitted to return to the configuration shown in FIG. 2C, and the gripping elements 46 and teeth 50 engage the connector 12, thus preventing movement between the connector 12 and the anterior anchor 40.

In an alternative embodiment, it is contemplated that the connector 12 can have notches 51, which interact with gripping elements 46 in a ratchet-and-pawl mechanism similar to that used in cable ties, providing a one-way adjustability, in which the posterior and anterior anchors can be moved toward each other, but not away from each other.

Figures 2E and 2F illustrate another embodiment of an anterior anchor 60 which is similar to the one illustrated in FIGs. 2C and 2D. In FIG. 2E, the gripping elements 62 and teeth 64 are oriented so that the anterior anchor can be deformed such that the top surface of the anchor is folded inward as illustrated in FIG. 2F. This is in contrast to the embodiment illustrated in FIG. 2D where the bottom surface of the anchor is folded inward. The teeth 64 in FIG. 2E are angled toward the top surface of the anterior anchor and engage the connector 12 of FIG. 1A such that they prevent movement of the anterior anchor along the connector 12 in the direction of the arrow in FIG. 2E, which would increase the distance between the anterior anchor and posterior anchor (not shown).

Figure 2G is a perspective view of a preferred embodiment where the anterior anchor is inflatable. The anterior anchor has a hollow, inflatable disc-shaped body 65 with a hole or other passageway 66 substantially in the middle of the body. Two gripping elements 67 project into the center of the hole or other passageway, although there can be as few as one or more than two. The gripping elements can have teeth 68 angled toward the top surface of the anchor. Alternatively, in a preferred embodiment, the gripping elements are in the form of a rough surface rather than the protruding elements as shown in FIG. 2G. Such a surface, which may be a sandpaper-like surface, creates enough friction to prevent movement in either direction along the connector. Optionally, two hooks 69 are located on the top surface of the anterior anchor. Hooks 69 facilitate grasping by a surgical instrument during deployment of the anterior anchor in the patient as described below. Alternatively, rather than hooks, there can be one or more graspable protrusions on the

body. In yet another embodiment, there are no hooks or graspable protrusions, and the body of the anchor is grasped directly to manipulate the anchor. In another embodiment, protrusions 69 are magnetic or otherwise sticky in nature to facilitate attachment to a surgical instrument.

5 An inflation tube 63 is used to inflate and deflate the anterior anchor. This inflation tube may or may not have a valve. In one preferred embodiment, the anterior anchor is filled with gas or fluid through the inflation tube and the fluid is held inside the anchor through an external (e.g. stopcock) valve controlled by the operator. When the inflation tube is cut at the end of the procedure, the inflation line is crimped closed thereby locking
10 the inflating substance inside the anchor. Alternatively, the shears used to cut the inflation line can be metal and an electrocautery current can be applied through the shears and to the inflation line to weld it closed.

 Figures 2H and 2I are cross sections of the anterior anchor of FIG. 2G, taken along the line D-D in FIG. 2G. The disc-shaped body 65 is readily deformable when in its
15 reduced profile (i.e., uninflated) configuration as shown in FIG. 2I. The body can be inflated with a substance delivered through the inflation tube 63. When anchor body is inflated, the anchor assumes its deployed (i.e. inflated) configuration as shown in FIG. 2H with the connector 12 of FIG. 1A passing through the hole 66 in the body of the anchor. In the deployed configuration, the gripping elements 67 and teeth 68 engage the connector
20 12 with sufficient pressure to prevent movement of the anchor along the connector 12 in the direction of the arrow in FIG. 2H, which would increase the distance between the anterior anchor and posterior anchor (not shown). Alternatively, rather than defined gripping elements and teeth, the surface of body which defines the sides of the hole or other passageway 66 can be configured such that when the anchor body is inflated, the
25 sides of the hole or other passageway expand to substantially close off the hole or other passageway and limit movement of the anchor relative to the connector through friction between the connector and the anchor.

 In FIG. 2I, the anterior anchor 65 is in its reduced profile (i.e. uninflated) configuration with the connector 12 of FIG. 1A passing through the hole 66 in the body of
30 the anchor. When in this configuration, the anchor body is readily deformable and the gripping elements 67 and teeth 68 do not significantly engage the connector 12. This

allows movement of the anterior anchor 65 along the length of the connector 12 in the directions illustrated by the arrows in FIG. 2I. Once the anterior anchor is in the desired position along the connector 12, the anterior anchor is inflated by a filling substance delivered through the inflation tube 63, and the anchor assumes its deployed (i.e. inflated) configuration as shown in FIG. 2H; the gripping elements 67 and teeth 68 engage the connector 12, thus restricting movement of the anterior anchor 65 in one or both directions along the length of the connector 12. The filling substance can be a gas, liquid, or material which changes phase with time (i.e. it may harden, cure, polymerize, or become a gel with time).

Figure 3A illustrates another embodiment of an anterior anchor 70 consisting of two parts, an anchor body 72 and a readily deformable collar 74. The anchor body and collar have a central hole or other passageway (76 and 78 respectively) through which the connector can pass. Preferably, the anterior anchor body is made of a semi-rigid polymer which can be deformed into a folded configuration with a reduced profile as illustrated in FIG. 3B. Preferably, the readily deformable collar 74 is permanently deformable; i.e., once deformed, it does not return to its original shape. As illustrated by the arrow in FIG. 3B, both the collar 74 and anchor body 72 can move along the connector 12 of FIG. 1A. Once the anchor body 72 is in the desired position, the collar 74 is crushed, such that the collar 74 engages the connector 12 and can no longer move along the length of the connector 12. This prevents the anchor body 72 from moving along the length of the connector 12 in the direction of the arrow illustrated in FIG. 3C, which would increase the distance between the anterior anchor and posterior anchor (not shown). Figure 3D illustrates an alternative embodiment of the anterior anchor 80, where the anchor body 82 and deformable collar 84 are a single piece.

In a preferred embodiment, the anterior anchor is made from a biocompatible, radio- or magneto-opaque polymer, but it can be made from various kinds of suitable materials known to those of skill in the art including metals, metal alloys, plastics, natural materials or combinations thereof as disclosed above. The anterior anchor can be solid, or alternatively, can be porous, mesh-like, umbrella-like or lattice-like. In a preferred embodiment, the anterior anchor is porous, mesh-like, umbrella-like or lattice-like to encourage fibrous ingrowth such that it becomes permanently attached to the stomach wall.

Coatings can be added to the anchor, or a mesh material such as polypropylene can be fixed to the anchor surface, such that it touches the anterior stomach wall and encourages tissue ingrowth. In other embodiments, the anterior anchor is solid and treated to discourage tissue ingrowth with materials such as PTFE or FEP which are generally hydrophobic and non-reactive. In other embodiments, the anterior anchor has a xenograft or allograft material attached to the anchor. In a preferred embodiment, the anterior anchor is disc-shaped and substantially flat, but those of skill in the art will recognize that other embodiments are possible.

Surgical Instruments

Figure 4A illustrates one embodiment of a tissue grasping instrument 200. The tissue grasper has a tubular outer sleeve 210 to which a portion of a handle 212 is attached at the proximal end. As shown in more detail in the blow-up, FIG. 4A, disposed within the outer sleeve 210 is a tubular inner member 214 which has an outer diameter such that it can slide within the outer sleeve 210 in the longitudinal axis of the outer sleeve 210 but cannot move substantially transverse to the longitudinal axis of the outer sleeve 210. At the proximal end of the inner member, a second portion of a handle 216 is attached. At the distal end of the inner member is a pair of jaws 220 which is connected to the inner member at a hinge point 222. When the distal end of the inner member 214 is displaced from the inside of the outer sleeve 210 such that the hinge point 222 is outside the outer sleeve, the jaws 220 assume their open position as depicted in FIG. 4A. As the hinge point 222 is withdrawn into the outer sleeve 210, the outer sleeve forces the jaws 220 into their closed position, as illustrated in FIG. 4B. The opening and closing of the jaws 220 can be accomplished by manipulation of the handle portions 212 and 216.

The distal end of the grasping instrument 200 is configured to cut, puncture, or dilate tissue when the jaws 220 are in the closed position. In one embodiment shown in FIG. 4B, the jaws 220 have screw-thread-shaped protrusions 224 on the surface. By rotating the instrument as it passes through tissue, the protrusions 224 facilitate the penetration of tissue, similar to a corkscrew. In another embodiment illustrated in FIG. 4C, the instrument has jaws 226 that form a sharp tip 228 when closed. In yet another embodiment, the jaws form a blade which can cut through tissues when in the closed

position. One of skill in the art would recognize that the above configurations can be combined, or that other configurations are possible which facilitate the passage of the tip of the instrument through the wall of the stomach or other tissue.

It also should be realized to one skilled in the art that the closed end of the grasping device does not have to be the only instrument responsible for cutting through the tissue; the central lumen 230 of the device can be utilized to assist in tissue penetration. For example, a needle (e.g. a Verres needle) 232 can be passed through the lumen and the needle 232 can make the initial puncture through the tissue. The configuration of the distal end of the grasper is meant to be a tissue dilator and facilitator of the entry into the stomach after the needle makes the initial puncture. For safety, the needle can be retracted as the tissue grasper dilates the tissue.

In the embodiment of the tissue grasper 200 illustrated in FIG. 4A, the inner member 214 and outer sleeve 210 have a central tunnel 230 that extends the length of the tissue grasper. The tunnel 230 allows for the passage of an expanding means such as a needle 232, or other instrument or device such as the posterior or anterior anchor described above, through the length of the tissue grasper as shown in FIG. 4A. The central tunnel is also adapted such that a radially dilating sheath can be inserted through it. The diameter of the central lumen is preferably at least 4 mm, but can be at least 5, 6, 7, 8, 9, 10, 11, or 12 mm. In an alternative embodiment, the distal jaws can be configured to close through an electromechanical means or purely magnetic means such that the inner member is not necessary.

Figure 5A illustrates one embodiment of an anchor implantation instrument 250. The implantation instrument has a tubular outer sheath 252 which has a handle 254 attached. At the distal end, the outer sheath flairs out to an increased diameter 255 to accommodate the anterior anchor in its substantially folded position as illustrated in FIG. 5C. Within the outer sheath is an anchor grasping instrument 256 similar to the tissue grasping instrument of FIG. 4A, made up of a tubular middle sleeve 260 and a tubular inner member 264. The tubular middle sleeve 260 has an outer diameter such that it can slide within the outer sheath 252 in the longitudinal axis of the outer sheath 252 but cannot move substantially transverse to the longitudinal axis of the outer sheath 252.

The tubular middle sleeve 260 of the anchor grasping instrument has a portion of a handle 262 attached at the proximal end 261 of the instrument. Disposed within the middle sleeve 260 is a tubular inner member 264 which has an outer diameter such that it can slide within the middle sleeve 260 in the direction of the longitudinal axis of the middle sleeve 260 but cannot move substantially in transverse to the longitudinal axis of the middle sleeve 260. At the proximal end of the inner member, a second portion of a handle 266 is attached.

The distal tip 263 of the instrument is illustrated in more detail in FIGS. 5B and 5C, with the inclusion of the anterior anchor 40 of FIG. 2A and connector 12 of FIG. 1A. Figure 5C is a side section view taken along the line C-C of FIG. 5B. At the distal end 263 of the inner member 264 is a pair of hooking members 270 which are connected to the inner member at a hinge point 272. When the distal end of the inner member 264 is displaced from the inside of the middle sleeve 260 such that the hinge point 272 is outside the middle sleeve, the hooking members 270 assume their open position as depicted in FIG. 5B. As the hinge point 272 is withdrawn into the middle sleeve 260, the middle sleeve forces the hooking members 270 into a closed position, as illustrated in FIG. 5C. The opening and closing of the hooking members 270 can be accomplished by manipulation of the handle portions 262 and 266.

The instrument is designed such that the anterior anchor is easily manipulated. When the anterior anchor is in its substantially folded or compressed configuration as in Fig. 5C, the entire anterior fastener assembly can be manipulated along the longitudinal axis of the connector 12. Fig. 5C depicts the assembly as it would be introduced over the connector 12 and into the patient. The operator pulls the connector 12 toward the operator such that the posterior anchor is urged toward the anterior anchor. When in position, the operator deploys anterior anchor 40. To deploy anterior anchor 40, outer sheath 252 is pulled back toward the operator. Middle sleeve 260 is then withdrawn proximally toward the operator as well. Hooking members 270 tend to fan out as the middle sleeve is pulled back and will release hooks 52. Once deployed, anterior fastener 40 is now fixed in a longitudinal position along the connector 12.

If the surgeon wants to readjust the anterior anchor, connector 12 is manipulated so that the hooks 52 of the anterior anchor are brought into contact with hooking members

270; middle sleeve 260 is advanced distally from the operator, permitting hooking members 270 to engage the hooks 52; such contact is facilitated by pulling back (proximally) on the connector 12. By manipulating the middle sleeve 260 over the hooking members 270, the hooks 274 on the ends of the hooking members 270 can engage the hooks 52 on the anterior anchor 40. The outer sheath 252 is then slid over the anterior anchor 40 (or the anchor-middle sleeve complex is withdrawn into the outer sheath 252), until it is compressed into an undeployed configuration as shown in FIG. 5C. As described above, when the anterior anchor 40 is in a substantially compressed configuration, it can move along the length of the connector 12 in either direction.

In an embodiment where an inflatable anterior anchor such as the one illustrated in FIGs. 2G-2I is utilized, a standard laparoscopic grasping instrument (with teeth) can be used to manipulate the anterior anchor. When the inflatable anterior anchor is in the uninflated position, it is sufficiently compliant such that it can easily be passed through a laparoscopic port prior to inflation and deployment or after it has been deflated for readjustment; the middle sheath may not be necessary because the compliance of the balloon enables easy compression into the outer sheath. The inflation tube 63 passes through the laparoscopic port and out of the patient. This allows the inflation tube 63 of the anchor to be temporarily opened or closed outside the patient allowing for deflation and reinflation until the anchor is in place. The inflation tube is then sealed and cut off, preferably substantially flush to the surface of the anterior anchor.

METHODS

Implantation of the Transgastric Fastening Assembly

Figure 6A depicts the initial step of a preferred embodiment of a surgical method to implant the transgastric fastening assembly. The first part of the procedure involves entering the stomach with an endoscope 300 and insufflating the stomach with a gas. When insufflated, the anterior wall of the stomach 302 is pushed toward the anterior abdominal wall 304 to create a potential space. After insufflation of the stomach, an incision is made in the skin and a standard laparoscopic port 306 is placed through the anterior abdominal wall 304 to a position wherein the distal end is in the potential space between the

abdominal wall 304 and the anterior wall of stomach 302. The laparoscopic port can be a radially dilating type port or similar port known in the art.

A particularly advantageous port is one which allows visualization of the individual abdominal layers as it is being pushed through the abdominal wall (well known to those skilled in the art). Use of such a port allows the surgeon to "see" the different layers of the abdominal wall from within the trocar (using a standard laparoscopic camera) as the trocar is advanced through the abdominal wall. The endoscopic light inside the stomach will be "seen" by the surgeon as the port approaches the inner layers of the abdominal wall because the endoscopic light source transilluminates through the layers of the stomach wall and inner layers of the abdominal wall. Such visualization is advantageous if the patient has a very thick abdominal wall (e.g. in a morbidly obese patient) because the surgeon needs to ensure that another organ (e.g. the colon) is not draped between the stomach and the posterior wall of the abdomen.

The tissue grasping instrument 200 of FIG. 4A is inserted through the port 306 with the jaws 220 in the closed position (with or without a needle projecting in front of the instrument) and is passed through the anterior wall of the stomach 302. When the jaws of the instrument are closed, the jaws define a sharp, dilating, and/or cutting configuration which can more easily advance through the stomach wall.

Figure 6B depicts the next step in a preferred method. The jaws of instrument 200 are used to grasp the posterior wall of the stomach 314. The posterior wall of the stomach 314 is lifted away from the retroperitoneum 316, allowing for access to the potential space of the lesser peritoneal sac 320. A needle 232, such as a Veress needle (well-known in the art, a Veress needle allows for easy and safe access into and between two serosal layers), is inserted through the central channel 230 of the instrument and passed through the posterior wall of the stomach 314 into the potential space of the lesser peritoneal sac 320. The potential space of the lesser peritoneal sac 320 is expanded by injection of a gas, such as carbon dioxide, through the needle 232. In other embodiments, the potential space is expanded using a liquid, gel, or foam. Alternatively, the space can be expanded using a balloon or other space expanding or space filling device; alternatively, a surgical instrument (e.g. electrocautery and/or blunt ended grasper, etc.) can be used in place of a needle to access the lesser peritoneum or to expand the potential space of the retroperitoneum 320.

Preferably, the expanded space of the lesser peritoneal sac can extend from the angle of His at the gastroesophageal junction to the pylorus.

In an alternative embodiment, the space is not expanded before the posterior anchor is placed. For example, in an embodiment where an inflatable posterior anchor is used, the potential space can be expanded by the anchor itself as it is inflated to its deployed configuration.

Figure 6C depicts the next step in a preferred embodiment. With a direct path from outside the patient to the lesser peritoneal sac 322, the needle 232 is withdrawn from the instrument 200. An optional dilation step can be performed at this stage in the procedure using a device such as a radially dilating sheath (e.g. InnerDyne STEPTM system; Sunnyvale, CA) inserted through the central channel 230 of the instrument. The dilating device expands the opening in the posterior wall of the stomach in such a way that the opening contracts down to a lesser profile after dilation. A posterior anchor 324 and connector 326, such as those depicted in FIGs. 1B, 1E or preferably 1F, in its reduced profile configuration, is passed through the central channel 230 of the instrument, through the posterior wall of the stomach 314, and deployed in the lesser peritoneal sac 322 as shown in FIG. 6C. Where the optional dilation step is performed, the posterior anchor 324 is passed through the dilating sheath. The connector 326 is preferably of sufficient length to pass from inside the lesser peritoneal sac 322 through the central channel 230 of the instrument and out of the patient's body. Figure 6D depicts the deployed posterior anchor 324 and connector 326 after the grasping instrument is withdrawn from the patient and tension is applied to connector 326 to pull the posterior anchor 324 against the posterior wall of the stomach 314.

Figure 7A illustrates the next step in the embodiment. The connector 326 is inserted through the hole or other passageway in an anterior anchor 40 of FIG. 5C, and the anchor implantation instrument 250 of FIGs. 5A, 5B and 5C is used to slide the anchor 40 through the laparoscopic port 306 into the abdomen of the patient. The anterior 302 and posterior 314 walls of the stomach are urged together, either by using the anchor implantation instrument 250 to urge the anterior wall 302 toward the posterior wall 314, or by pulling on the connector 326 and posterior anchor 324 to urge the posterior wall 302 of the stomach toward the anterior wall 314, or by a combination of the two methods. Once

the anterior anchor 40 is in the desired position, the anterior anchor 40 is placed in its deployed configuration by manipulating the anchor implantation instrument 250 as described above.

In a preferred embodiment, the inflatable anterior anchor of FIGs. 2G-2I is used, and the use of the implantation instrument of FIG. 5C is optional. After the anterior anchor is in the desired position, the anterior anchor is inflated with a filling substance through the inflation tube until it is in its deployed configuration. The gripping elements 67 and teeth 68 are thus engaged against the connector 326. The anchor implantation device 250 can then be withdrawn from the patient's abdomen.

With the transgastric fastening assembly complete, the surgeon can examine the resulting configuration of the stomach using an endoscope. If the anterior anchor is not in the desired location, its placement along the connector can be adjusted as described above. Alternatively, in another embodiment, the anterior anchor can be urged closer to the posterior anchor simply by pushing it along the connector without using the implantation device to capture the anchor and deform it into its reduced profile configuration.

In the preferred embodiment, the anterior anchor can be deflated, allowing the anterior anchor to be repositioned, and then reinflated to engage the connector. Figure 7B illustrates the transgastric fastening assembly with the anterior anchor 40 in its deployed configuration on the connector 326 and the anchor implantation instrument removed from the patient's abdomen. The anterior 302 and posterior walls 314 of the stomach have been urged closer together by the transgastric fastening assembly. Whether the walls of the stomach are urged into contact or not is determined by the surgeon.

Figure 7C depicts the transgastric fastening assembly in its final configuration after deployment. Once the surgeon is satisfied that the transgastric fastening assembly is properly placed, a cutting implement, well-known to those of skill in the art, is inserted through the laparoscopic port and the connector 326 is cut, preferably flush to the anterior anchor 40. In the preferred embodiment, where inflatable anchors are used, the hollow connector and inflation tube are sealed prior to, or as a result of, cutting, preventing anchor deflation. Alternatively, if a filling substance which hardens with time is used, it may not be necessary to seal the connector or inflation tube prior to cutting if the filling substance is sufficiently hard or viscous such that it will not leak from the connector or inflation tube.

When more than one transgastric fastening assembly is to be implanted, it is preferred to insert all of the posterior anchors and connectors before attaching any anterior anchors. This is in contrast to attempting to place one complete transgastric fastening assembly and then subsequent assemblies. While possible, if one were to place entire
5 fastening assemblies in series, each successive assembly would be more difficult to place because the volume of the stomach would be progressively reduced resulting in more difficult visualization each time.

Figure 8A depicts an embodiment in which two posterior anchors 330 and connectors 332 are deployed in the expanded lesser peritoneal sac. In this embodiment,
10 there is one laparoscopic port 334 for each connector 332. Alternatively, there may be more fasteners placed than incisions and laparoscopic ports. Depending on how far apart the fasteners are placed, a given laparoscopic port can be used to implant a plurality of transgastric implants. This can be accomplished because there is significant mobility of the stomach and/or abdominal wall which allows for different points along the anterior wall of
15 the stomach to be accessed without having to create another hole through the abdominal wall.

When it is desired to place more than one set of transgastric assemblies and in particular when the assemblies are placed concurrently rather than sequentially, the surgeon is afforded the opportunity to test (e.g measuring stomach volume, resistance to flow,
20 assessing mucosal integrity, etc.) varying tensions on one or more of the fastening assemblies, all under endoscopic inspection. After the desired number of posterior anchors and connectors are deployed in the patient, the configuration of the stomach can be tested by applying tension to the connectors. Figure 8B depicts temporary clamps 336 which sit on top of the ports 334. Connectors 332 can be pulled from outside the abdomen to urge
25 the posterior wall of the stomach 340 toward the anterior abdominal wall 342. One or more clamps 336 can then be closed to hold the stomach in a test position. To determine if the posterior anchors 330 are in the desired location, an endoscope 344 can be used to view the configuration and the tension that the stomach will endure after the anterior anchors are placed.

30 In an alternative embodiment, the stomach is fastened to the abdominal wall rather than there being a free space between the anterior gastric wall and the peritoneum of the

abdominal wall. The initial steps are as discussed above. After the posterior anchors are placed, their position can be tested as depicted in FIG.8B to simulate the configuration after the anterior fastener is placed. Next, the outer laparoscopic port is pulled back so that the anchor deploying instrument directly contacts and sits within the tissues of the muscular abdominal wall. Once the outer laparoscopic port is pulled back, the anterior anchor can be deployed within the abdominal wall musculature and the connector can be cut flush with the anterior fastener. In an embodiment where the inflatable anterior anchor is used, after the anterior anchor is deployed within the abdominal wall musculature, the inflation tube is cut, preferably flush with the anterior anchor.

Reversal of the Gastric Volume Reduction Procedure

The connector of a preferred embodiment of the deployed transgastric fastening assembly, as illustrated in FIG. 7C, can be cut at a point between the anterior and posterior anchors, which results in reversal of the gastric volume reduction. The connector is preferably made to resist corrosion from stomach acid, but is able to be cut by a cutting implement advanced through an endoscope into the stomach. Materials suitable to prevent corrosion and yet allow cutting include plastics such as polyurethane, silicone elastomer, polypropylene, PTFE, PVDF, or polyester, metals and metal alloys such as stainless steel, nickel-titanium, titanium, cobalt-chromium, etc. Once the connector is cut, the walls of the stomach are free to move away from one another, thereby reversing the procedure. Reversal of the procedure can occur at any time (days to years) after the procedure. In a preferred embodiment, the anchors remain in the gastric wall permanently even after the connector is cut or otherwise divided. Alternatively, the anchors can in part or in whole be manufactured from a bioabsorbable material such that the anchors will eventually be absorbed by the body. In the case of bioabsorbable anchors, it is preferable to have a connector which is at least in part bioabsorbable. In another embodiment, substantially all of the elements of the transgastric fastening assembly are made of bioabsorbable materials, with the intent that over the desired period of time, the entire assembly will be absorbed by the body, reversing the procedure without any additional actions required by a doctor.

Even if there is some degree of fusion between the mucosa around the connector at the region of the assembly, once the connector is cut or absorbed, the walls will tend to

move apart over time. Alternatively, a balloon or other dissection device is introduced through an endoscope and used to separate the walls of the stomach at the point of fusion.

Treatment of Disease Conditions

- 5 The devices, methods and instruments disclosed above can be used to treat obesity and other diseases involving the gastrointestinal tract, such as gastroesophageal reflux disease (GERD). Figure 9 depicts three transgastric fastening assemblies 400 deployed in the stomach. The dashed lines represent boundaries of the divisions of the stomach: the cardia of the stomach 402, the fundus of the stomach 404, the body of the stomach 406, 10 the antrum of the stomach 408, and the pyloric sphincter 410. In a preferred embodiment, the fastening assemblies are not implanted in the antrum 408 in order to maintain the normal digestion process of the stomach. Normal digestion occurs in the antrum which precedes passage of food into the duodenum. In stopping short of the antrum 408, the implants replicate the degree of volume reduction of the M&M procedure.
- 15 Food ingested by the patient follows a physiologic pathway for digestion depicted by the arrow in FIG. 9. It travels through the esophagus 412 and enters the cardia of the stomach 402. The food is digested in the stomach and pushed toward the duodenum 414 as chyme for further digestion. The preserved antrum 408 will allow for relatively physiologic digestion and emptying into the duodenum 414 akin to the M&M procedure.
- 20 With transgastric fastening assemblies 400 in place, food which leaves the esophagus 412 and enters the stomach, results in increased wall tension on the lesser curvature of the stomach 416 as the greater curvature of the stomach 418 will be restricted from the food pathway. The path of least resistance will be the path toward the pylorus 410 and duodenum 414. The increased wall tension of the stomach will result in a feeling of satiety 25 in the patient, leading to decreased food intake and weight loss. Although three assemblies are shown in FIG. 9, there may be as few as one or as many as ten depending on the degree of volume reduction desired. Such flexibility in number of devices as well as the ability of the surgeon to tune the tension between the anterior and posterior fasteners is advantageous. Such flexibility may enable, for example, reversal of a few fasteners rather 30 than all the fasteners, such that the volume reduction procedure is partially reversed.

In another embodiment, a transgastric fastening assembly is placed in the antrum 408 or the region just proximal to the pyloric sphincter 410 if deemed necessary by the gastroenterologist and/or surgeon. Such a configuration would not reduce the volume of the stomach but would cause a feeling of fullness similar to a gastric outlet obstruction, leading to decreased food intake and weight loss. The fasteners in this region can also conduct a current to electrically stimulate the stomach to simulate satiety.

In another embodiment, a transgastric fastening assembly may be required at the region of the cardia 402 to treat morbid obesity in a similar manner to that utilized with the LAP-BAND™. In this embodiment, the transgastric fastening assembly is not utilized to reduce the volume of the stomach, but to create a restriction to the inflow of food.

In another embodiment, the surgeon or gastroenterologist may choose to treat a disease such as gastroesophageal reflux disease (GERD) with a transgastric fastening assembly in the cardia region. Such a configuration would maintain the position of the GE junction in the abdomen and potentially great a barrier resistance to reflux contents.

In another embodiment, the disclosed method in combination with the transgastric fastening assemblies can be adapted to attach a gastrointestinal organ to the abdominal wall which in addition to reducing volume can also create a kink in the organ. The kink would cause a resistance barrier (in addition to volume reduction) to gastrointestinal contents, and can be useful to treat reflux disease or morbid obesity.

Such a kink would also fix the gastrointestinal region to the abdominal wall can also maintain the reduction of a hiatal hernia in the abdominal compartment (e.g. in reflux disease). A major component of reflux disease is a hiatal hernia in which the gastroesophageal junction freely slides from the abdomen to the mediastinum. A percutaneously placed suture or anchor in the region of the gastric cardia and/or fundus can tether the junction to the abdominal wall and confine the junction to the abdomen.

In other embodiments, the devices and methods of this invention can assist in the implantation of devices such as stents, meshes, stitches, or tubes in the gastrointestinal tract. The major technical difficulty encountered in placing stents, tubes, and meshes inside the lumen of the gastrointestinal tract is that they tend to migrate because the walls of such devices do not adhere to slippery mucosa. A transgastric or transintestinal fastener, implanted with the current instrumentation would solve this problem. Such a method

would be particularly useful in the attachment of the stent part of the stent-sleeve system outlined in patent application WO 04049982 , or the mesh of patent application WO03086247A1. In another example, devices such as those disclosed in patent US6773441 attempt to place an endoscopic stitch to tether the cardia of the stomach to the fundus to treat reflux disease. Such stitches are tenuous in the long term because they do not necessarily penetrate the serosa. Even if the stitches penetrate the serosa, they tend to erode through the wall with time because of their thin profile and an inability of the endoscopic operator to control tension on the suture when it is placed. With the methods and devices of this invention, such an endoscopic suture can be buttressed with a percutaneously placed fastener.

Other Uses for the Disclosed Devices, Instruments, and Methods

Although the described methods are focused on the implantation of transgastric fastening assemblies to reduce the volume of the stomach, the methods and devices can easily be expanded to the percutaneous placement of other types of devices such as neurostimulators, gastric muscle stimulators, gastric balloons, bulking devices inside the wall of a gastrointestinal organ, devices placed in the lesser peritoneal sac along the autonomic nerve plexus, along the vagus nerve, on parts of the diaphragm, or placed on or along the pancreas.

In some embodiments, other devices are implanted using the described methods to place devices inside or outside the stomach; inside or outside the lesser sac of the peritoneum; inside or beside a structure within the retroperitoneum; inside, beside, or outside the duodenum, pylorus, or gastroesophageal junction. Implanted devices include but are not limited to the anchor devices and transgastric fastening assemblies described above, neuromodulators, direct muscle stimulators, stents, meshes, stent-grafts, stitches, and bulk forming agents.

In one such embodiment, a transgastric fastening assembly serves to reduce the volume of the stomach as well as provide for electrical stimulation. In this embodiment, an electrical signal runs through electrodes in the transgastric fastener assembly to alter the contraction patterns of the stomach or to electrically stimulate a feeling of satiety as well as reduce the volume of the stomach. Thus, fastener assemblies of the present invention can

become electrodes which are useful, for example, for gastric electrical stimulation. Methods and devices of this invention can also be used to place sutures in the stomach or pylorus to treat reflux disease. Such suturing would be facilitated by the placement of multiple ports through the walls of the stomach; this would be highly beneficial over current fully
5 endoscopic methods of placing sutures. Any of these methods and devices could be used in combination with or in place of the transgastric fastening assemblies to induce weight loss in a patient.

In other embodiments, the novel methods, implantation devices, and fasteners of this invention are used to implant devices in one wall of a gastrointestinal organ without
10 volume reduction. In one example of such an embodiment, illustrated in FIGs. 10A and 10B, a balloon-like device is deployed in the stomach to displace volume rather than to reduce volume from the outside. The balloon 430 is the equivalent of the posterior anchors in the above embodiments. In this embodiment, after initial insufflation of the stomach and placement of a laparoscopic port 306 in the abdominal wall 304, an instrument is used to
15 penetrate only the anterior wall of the stomach 302 and place an inflatable intragastric balloon 430. Inflation is achieved through connector 432 and the balloon is placed within the interior of the stomach 428, as illustrated in FIG. 10A. When inflated, the balloon 430 is preferably spherical in shape such that it occupies a portion of the stomach volume when inflated. In the embodiment shown, the connector also acts as the inflation tube for
20 inflating the intragastric balloon. In another embodiment, there is a separate inflation tube in addition to the connector. As discussed above, a valve can be located between the anchor and the connector, or alternatively outside the patient. Preferably after the intragastric balloon is inflated, an anterior anchor 434 is deployed on the connector 432 as described previously, the connector is cut, preferably flush with the anterior anchor, and
25 the laparoscopic port is removed, as shown in FIG. 10B. In the preferred embodiment where an inflatable anterior anchor 434 is used, the inflation tube is also cut, preferably flush with the anterior anchor. Other devices which are only implanted in the anterior gastric wall include neurostimulators, muscular stimulators, sensors, pharmaceutical delivery devices

30 In another embodiment, an extragastric balloon is used to reduce the volume of the stomach and/or create a barrier to the flow of food and a restrictor to the flow of food.

Figure 11a depicts the balloon 430 in the undeployed configuration. The balloon is placed through trocar port 306 which is first placed as described above. Figure 11B shows an embodiment of an extragastric balloon 430 in its deployed state. The balloon 430 is attached to the abdominal wall by any of the percutaneous fastener-connector assemblies and methods described above. Stem 432 is the residual from the connector used to place the balloon with an optional access port for further inflation and/or deflation after the balloon is placed. In some embodiments (Fig.11c), the posterior portion of the balloon 436 is fixed to the outer portion or inner portion of the stomach using any of the fastening systems described above. The posterior portion of the balloon can also be fixed to the anterior gastric wall with a fastener delivered through the stomach with an endoscope. The extragastric balloon can be placed anywhere along the stomach, even at the region 1-5cm below the gastroesophageal junction at the same place where laparoscopically placed gastric bands are currently placed.

15 Alternative Implantation Techniques

Although the methods described in this disclosure focus on an entirely percutaneous surgical procedure where general anesthesia, paralysis, and pneumoperitoneum are not used, it is conceivable that the surgeon may want to use general anesthesia, pneumoperitoneum, and paralysis. Such may be the case when the adhesions behind the stomach are severe and it will be safer to perform a standard laparoscopic procedure to implant the devices of this invention. Alternatively, it may be simple surgeon preference and comfort level which dictates whether or not general anesthesia and/or pneumoperitoneum are used. In such a case, the percutaneous instruments and implants described in this invention will be used, but the surgeon will additionally have the advantage of laparoscopic instrumentation and tools which can free the posterior gastric wall from the retroperitoneum under direct visualization.

It is also possible that a part of, or the entire procedure, is performed with a fluoroscope, MRI, CAT scan, or ultrasound, in which the structures behind the stomach are can be visualized.

In one embodiment, the fastening system is adapted for placement during laparoscopy, Fig. 12. In this case, a laparoscopic tool is provided which has a reversibly attached fastener. Grips 520 reversibly grip fastener 510. Any of a variety of gripping mechanisms can be employed to retain the fastener 510 on laparoscopic tool 500.

- 5 Connector 332 is substantially similar to any of the connectors described above except that the posterior fastener 510 is not attached to connector 332 when it is inserted through the anterior abdominal wall. The surgeon places laparoscopic tool 500 behind the stomach 428 of the patient and connector 332 is advanced through lumen 545 formed in patient's skin 535 and anterior abdominal wall 530. Connector 332 is then further advanced
- 10 percutaneously through first and second walls 540 and 542 of stomach 428. In one embodiment, the connector and lumen 545 are one unit in which the connector can be controlled independently of the lumen 545.

- When the connector 332 reaches the posterior fastener 510, gripping elements 520 are released by the surgeon through a mechanism which is integrated into the laparoscopic tool 500. Connector 332 is fixed to posterior fastener 510 through a locking mechanism.
- 15 Mechanisms of locking connector 332 to posterior fastener 510 are well-known to those skilled in the art of mechanical fixturing. Some or all of the fixturing mechanisms may reside on the connector or on the fastener. In another embodiment, the gripping force of the grippers 520 can be overcome by force applied by the surgeon on connector 332.
- 20 Mechanisms of locking other than mechanical also exist and include magnetic, electromagnetic, and adhesive means.

- An anterior fastener 550 (Fig. 13) is then placed over the connector shown in Figure 12 by the methodology and devices described above. The walls of the stomach are urged together to create a resistance to the flow of food within the stomach (Fig. 14). 570 depicts
- 25 one side of the stomach after the walls of the stomach are urged together. It is this side of the stomach where food enters. Its volume and capacity are now reduced as compared to its original volume and capacity. Although not shown, connector 332 is truncated at the level of the anterior fastener by any of the mechanisms described and depicted above.

- Although the present invention has been described in the context of certain
- 30 preferred or illustrative embodiments, it should be understood that the scope of the

exclusive right granted by this patent is not limited to those embodiments, but instead is the full lawful scope of the appended claims.

WHAT IS CLAIMED IS:

1. A method for reducing the interior volume of an organ, comprising:
passing a first end of a surgical instrument through a patient's skin, through a first exterior surface of the organ, through the interior of the organ, and thence through a second exterior surface of the organ, so that the surgical instrument traverses the organ;
deploying a first anchor from the surgical instrument wherein the first anchor is located adjacent to the second exterior surface of the organ;
partially or completely withdrawing the surgical instrument;
deploying a second anchor wherein the second anchor is located adjacent to the first exterior surface of the organ; and
providing a connector between the first and second anchors, wherein the length of the connector between the first and second anchors is such that the first and second anchors urge the first and second exterior surfaces of the organ toward each other, thereby reducing the volume of the organ.
2. The method of Claim 1, wherein said first and second anchors are deployed from the same surgical instrument.
3. A method of reversing the volume reducing procedure of Claim 1 comprising:
providing a patient having one or more connectors in an organ, wherein the connectors have been implanted utilizing the method of Claim 1; and
cutting or otherwise dividing said one or more connectors between the first and second anchors.
4. The method of Claim 1, wherein the organ is a gastrointestinal organ.
5. The method of Claim 1, further comprising creating a space outside the organ adjacent to the second exterior surface thereof by introducing a volume-filling substance into a potential space adjoining said second exterior surface.
6. The method of Claim 5, wherein said potential space is expanded by the injection of a gas, liquid, gel, foam, or solid, into the potential space.
7. The method of Claim 5, wherein said potential space is expanded by the inflation of a balloon placed in the potential space.

8. The method of Claim 5, wherein said potential space is expanded by blunt dissection.

9. The method of Claim 4, further comprising insufflating the organ prior to said passing of said first end of the surgical instrument through the first exterior surface of the organ.

10. The method of Claim 5, wherein said patient's skin is overlying the patient's stomach, said organ is the patient's stomach, said first exterior surface is the anterior wall of the stomach, said second exterior surface is the posterior wall of the stomach, and said potential space is the lesser sac of the peritoneum.

11. The method of Claim 10, further comprising urging the anterior and posterior walls of the stomach closer together by shortening the length of the connector between the first and the second anchors.

12. The method of Claim 10, wherein the surgical instrument is inserted into the patient's abdomen by directly penetrating the patient's skin and abdominal wall, by passing the surgical instrument through a laparoscopic port, or by passing the surgical instrument through an incision in the patient's skin and abdominal wall.

13. A method for reducing the volume of a patient's stomach, comprising:
passing a first anchor through a patient's abdominal skin, and through the patient's anterior and posterior stomach walls, wherein the first anchor is in a reduced profile configuration;

deploying said first anchor within the patient;

passing a second anchor through the skin of a patient's abdominal wall;

deploying the second anchor within the patient;

connecting the first and second anchors by means of a connector that passes through the stomach;

urging the first and second anchors toward each other; and

holding the anterior and posterior walls of the stomach close together with said first and second anchors.

14. The method of Claim 13, wherein said second anchor is deployed within the peritoneal cavity.

15. The method of Claim 13, wherein said second anchor is deployed between the abdominal skin and the outermost peritoneum of the abdominal cavity.

16. The method of Claim 13, wherein passing said second anchor through the patient's abdominal skin is done while said second anchor is in a reduced profile configuration, further comprising said second anchor subsequently being in a deployed configuration.

17. The method of Claim 13, wherein said reduced profile configuration of said first anchor is substantially folded, and/or compressed, and/or uninflated, and said deployed configuration is substantially unfolded and/or uncompressed and/or inflated.

18. The method of Claim 16, wherein said reduced profile of said second anchor is substantially folded, and/or compressed, and/or uninflated, and said deployed configuration is substantially unfolded and/or uncompressed and/or inflated.

19. The method of Claim 14, wherein mucosa of said anterior and posterior walls of the stomach do not contact one another.

20. The method of Claim 14, wherein mucosa of said anterior and posterior walls contact one another.

21. The method of Claim 14, wherein said holding step is performed by adjusting the relative position of the connector and the second anchor, and then configuring the second anchor to engage the connector in a manner that prevents a change in the distance between the first and second anchors.

22. The method of Claim 13, further comprising delivering an electrical signal to said anterior stomach wall through said second anchor.

23. The method of Claim 13, further comprising delivering an electrical signal to said posterior stomach wall through said first anchor.

24. The method of Claim 14, wherein the method is performed without the use of a general pneumoperitoneum;

and wherein an endoscope is not used for mechanical manipulation.

25. The method of Claim 14, further comprising imaging of the patient's abdomen using fluoroscope, MRI, CAT scan, or ultrasound, during insertion of either of said first or second anchors into the patient's abdomen.

26. A fastening assembly, comprising:

a first anchor, a second anchor, and a connector;

wherein said first anchor comprises a relatively planar body attached to said connector;

said body of said first anchor having a relatively planar deployed profile and a reduced profile configuration;

wherein said second anchor comprises a relatively planar body, a hole or other passageway approximately in the center of the body of sufficient diameter to allow passage of the connector through said hole or other passageway, and one or more gripping elements within said hole or other passageway;

said body of said second anchor having a relatively planar deployed profile and a reduced profile configuration; and

wherein said gripping elements prevent movement of said second anchor along the longitudinal axis of the connector in the direction away from said first anchor when the connector is disposed in said hole or other passageway and when said second anchor is in its deployed configuration.

27. The fastening assembly of Claim 26, wherein said body has two or more attachment structures accessible from a top surface of said body.

28. The fastening assembly of Claim 26, wherein said reduced profile configuration is substantially folded, and/or compressed, and/or uninflated, and said deployed configuration is substantially unfolded, and/or uncompressed, and/or inflated.

29. The fastening assembly of Claim 28, wherein said gripping elements do not prevent movement of said second anchor along the longitudinal axis in either direction when said connector is disposed in said hole or other passageway and when said second anchor is in its reduced profile configuration.

30. The fastening assembly of Claim 26, wherein said body of said first anchor is inflatable;

wherein said first anchor has a relatively planar deployed profile when inflated and a reduced profile configuration when uninflated;

wherein said first anchor is readily deformable when uninflated; and

wherein said connector is hollow and configured to allow inflation of said first anchor by delivery of a filling substance through said connector to said first anchor.

31. The fastening assembly of Claim 26, wherein said body of said second anchor is inflatable;

wherein said second anchor has a relatively planar deployed profile when inflated and a reduced profile configuration when uninflated; and

wherein said second anchor is readily deformable when uninflated.

32. The fastening assembly of Claim 31, wherein said gripping elements do not prevent movement of said second anchor along the longitudinal axis in either direction when said connector is disposed in said hole or other passageway when said second anchor is in its reduced profile configuration.

33. The fastening assembly of Claim 31, wherein said second anchor has an inflation tube connected to said second anchor body configured to allow inflation of said second anchor by delivery of a filling substance through said inflation tube to said second anchor body.

34. The fastening assembly of Claim 30, wherein said filling substance hardens, cures, polymerizes, or become a gel over time.

35. The fastening assembly of Claim 33, wherein said filling substance hardens, cures, polymerizes, or become a gel over time.

36. The fastening assembly of Claim 26, wherein said first anchor, said second anchor, or both said first and said second anchors, can deliver an electrical signal to tissue when placed in contact with said tissue.

37. A surgical instrument comprising:

a proximal end, a distal end, and a shaft with a channel;

said distal end comprising a tissue penetrator adapted to penetrate a wall of a hollow organ, and a tissue grasper adapted to grasp a wall of a hollow organ;

an expander within the channel of the surgical instrument;

wherein the expander can be deployed from within the channel of the instrument and extend through a wall of the hollow organ while the wall is being grasped by the tissue grasper, and is adapted to introduce a volume-filling material or device beyond the hollow organ.

38. A surgical instrument comprising:

a proximal end, a distal end, and a channel;

wherein said distal end comprises a tip which has an open configuration and a closed configuration;

wherein in said closed configuration, said tip is adapted to penetrate a wall of a hollow organ;

wherein in said open configuration, said tip can grasp the wall of the hollow organ.

39. The surgical instrument of Claim 38, wherein said channel comprises a diameter of about 2 mm to 12 mm.

40. An anchor implantation instrument comprising:

an outer sheath, a middle sleeve disposed within the outer sheath, an inner member disposed within the middle sleeve, and a channel within the inner member;

wherein said middle sleeve has an outer diameter and a longitudinal axis, and the outer diameter of said middle sleeve is such that it can slide within the outer sheath along the longitudinal axis of the middle sleeve;

wherein the inner member has an outer diameter, and the outer diameter of said inner member is such that it can slide within the middle sleeve along the longitudinal axis of the middle sleeve; and

wherein said inner member has a grasper suitable for grasping a foldable anchor and wherein said outer sheath has a distal portion which has an inner diameter sufficient to hold a foldable anchor in its folded configuration.

41. A biocompatible surgical anchor adapted for use in connecting the walls of a hollow organ or anchoring a device to the wall of a hollow organ, comprising:

a body portion, wherein said body portion is foldable and can assume a relatively planar deployed configuration and a reduced-profile folded configuration, wherein the body portion has an opening extending therethrough; and

a gripper on the body portion adapted to grip an elongated connector as such a connector extends through the opening, wherein the gripper engages such a connector when the anchor is in the deployed configuration, and releases such a connector when the anchor is in the folded configuration.

42. The surgical anchor of Claim 41, wherein the gripper, when engaging a connector, allows movement of the connector in one direction relative to the anchor but prevents movement of the connector in an opposite direction relative to the anchor.

43. A biocompatible surgical anchor adapted for use in connecting the walls of a hollow organ, comprising:

a body portion, wherein said body portion is inflatable and can assume a relatively planar deployed configuration when inflated and a reduced-profile configuration when uninflated, wherein the body portion has an opening extending therethrough; and

a gripper on the body portion adapted to grip an elongated connector as such a connector extends through the opening, wherein the gripper engages such a connector when the anchor is inflated, and releases such a connector when the anchor is uninflated.

44. The surgical anchor of Claim 43, wherein the gripper, when engaging a connector, allows movement of the connector in one direction relative to the anchor but prevents movement of the connector in an opposite direction relative to the anchor.

45. The surgical anchor of Claim 43, wherein said anchor has an inflation tube connected to said body portion configured to allow inflation of said body portion by delivery of a filling substance through said inflation tube to said body portion.

46. The surgical anchor of Claim 45, wherein said filling substance hardens, cures, polymerizes, or become a gel over time.

47. The surgical anchor of Claim 43, wherein said anchor has one or more attachment structures accessible from a top surface of said body.

48. A fastening assembly, comprising:

a first anchor, a second anchor, and a connector;

wherein said first anchor comprises an inflatable body attached to said connector;

said body of said first anchor having a deployed configuration when inflated that is relatively spherical and a reduced profile configuration when uninflated wherein it is readily deformable;

wherein said connector is hollow and configured to allow inflation of said first anchor by delivery of a filling substance through said connector to said first anchor;

wherein said second anchor comprises a relatively planar body, a hole or other passageway approximately in the center of the body of sufficient diameter to allow passage of the connector through said hole or other passageway, one or more gripping elements projecting into said hole or other passageway, and one or more attachment structures accessible from a top surface of said body;

said body of said second anchor having a relatively planar deployed profile and a reduced profile configuration wherein it is substantially folded; and

wherein said gripping elements prevent movement of said second anchor along the longitudinal axis of the connector in the direction away from said first anchor when the connector is disposed in said hole or other passageway when said second anchor is in its deployed configuration.

49. The fastening assembly of Claim 48, wherein said gripping elements do not prevent movement of said second anchor along the longitudinal axis in either direction when said connector is disposed in said hole or other passageway when said second anchor is in its reduced profile configuration.

50. The fastening assembly of Claim 48, wherein said filling substance hardens, cures, polymerizes, or become a gel over time.

51. The fastening assembly of Claim 48, wherein said body of said second anchor is inflatable;

wherein said second anchor has a relatively planar deployed profile when inflated and a reduced profile configuration when uninflated; and

wherein said second anchor is readily deformable when uninflated.

52. The fastening assembly of Claim 51, wherein said gripping elements do not prevent movement of said second anchor along the longitudinal axis in either direction when said connector is disposed in said hole or other passageway when said second anchor is in its reduced profile configuration.

53. The fastening assembly of Claim 52, wherein said second anchor has an inflation tube connected to said second anchor body configured to allow inflation of said second anchor by delivery of a filling substance through said inflation tube to said first anchor body.

54. The fastening assembly of Claim 53, wherein said filling substance hardens, cures, polymerizes, or become a gel over time.

55. A method of fastening a device to one wall of a gastrointestinal organ comprising;

passing said device through a patient's abdominal skin, and through the one wall of the gastrointestinal organ, wherein the device is in an undeployed configuration;

passing an anchor through the patient's abdominal skin wherein said anchor is in an undeployed configuration;

deploying said device;

connecting said device and said anchor by means of a connector that passes through the wall of said gastrointestinal organ;

wherein said connector prevents said device and second anchor from moving apart; and deploying said anchor to its deployed configuration.

56. The method of Claim 55 wherein said device comprises an inflatable body attached to said connector;

said body of said device having a deployed configuration when inflated that is relatively spherical and a reduced profile configuration when uninflated wherein it is readily deformable;

wherein said connector is hollow and configured to allow inflation of said device by delivery of a filling substance through said connector to said device;

wherein said anchor comprises a relatively planar body, a hole or other passageway approximately in the center of the body of sufficient diameter to allow passage of the connector through said hole or other passageway, one or more gripping elements associated with said hole or other passageway, and one or more attachment structures accessible from a top surface of said body;

said body of said second anchor having a relatively planar deployed profile and a reduced profile configuration wherein it is substantially compressed; and

wherein said gripping elements prevent movement of said anchor along the longitudinal axis of the connector in the either direction away from said device when the connector is disposed in said hole or other passageway and when said anchor is in its deployed configuration.

57. The method of Claim 56, wherein said filling substance hardens, cures, polymerizes, or become a gel over time

58. The method of Claim 56, wherein said body of said anchor is inflatable; wherein said second anchor has a relatively planar deployed profile when inflated and a reduced profile configuration when uninflated; and

wherein said second anchor is readily deformable when uninflated.

59. The method of Claim 58, wherein said gripping elements do not prevent movement of said second anchor along the longitudinal axis in either direction when said connector is disposed in said hole or other passageway when said second anchor is in its reduced profile configuration.

60. The method of Claim 59, wherein said anchor has an inflation tube connected to said anchor body configured to allow inflation of said anchor by delivery of a filling substance through said inflation tube to said anchor body.

61. The method of Claim 60, wherein said filling substance hardens, cures, polymerizes, or become a gel over time.

62. The method of Claim 55 wherein said device is an electrical stimulator.

63. The method of Claim 62 further comprising delivering an electrical signal to the wall of said gastrointestinal organ through said electrical stimulator.

64. The method of Claim 55 wherein said device is adapted to deliver a medicine.

65. The method of Claim 10, wherein mucosa of said anterior and posterior walls contact one another.

66. A surgical device system, comprising:
a laparoscopic instrument comprising a fastener adapted to receive a connector having a proximal end and a distal end; and
a grasping mechanism.

67. The surgical device system of claim 66, wherein said grasping mechanism is located on said instrument.

68. The surgical device system of claim 66 wherein the grasping mechanism is located on the connector.

69. The device of Claim 67 or 68, wherein the grasping mechanism comprises an actuator that is controllable at the proximal end of said connector or said instrument.

70. The device of Claim 69, wherein said actuator is adapted to activate the grasping mechanism to reversibly engage the fastener.

71. The device of Claim 66, wherein the connector further comprises a removable outer sheath.

72. A method of treating a patient, comprising:

positioning a first surgical device having a reversibly engaged first fastener near an organ of the patient;

positioning or forming an elongate member in the patient, comprising a proximal end, a distal end and a circumferential wall forming a lumen sized to receive a connector therethrough, wherein the connector has a proximal end and a distal end;

engaging the first fastener with the connector, the connector advanced through the lumen of the elongate member, the elongate member thence percutaneously applied through a first portion of the organ and thence through a second portion of the organ;

releasing the first fastener from the first surgical device such that the first fastener remains engaged with the connector;

advancing a second surgical device through the lumen, wherein the second surgical device comprises a second fastener configured to track the connector;

urging the first and second portions of the organ toward each other with the first and second fasteners;

releasing the second fastener from the second surgical instrument such that the second fastener engages the connector; and

cutting the connector such that the connector has a length that spans the first and second fasteners.

73. The method of Claim 72, wherein the organ is the patient's stomach.

74. The method of Claim 72, wherein urging the first and second portions of the organ reduces the volume of the organ and creates a resistance to the flow of material into the organ.

75. The method of Claim 72, wherein the elongate member is a laparoscopic port that is positioned in the patient.

76. The method of Claim 72, wherein the elongate member and the connector are one device.

77. The method of Claim 73, wherein the urging causes the mucosa of the first and second portions of the patient's stomach to contact each other.

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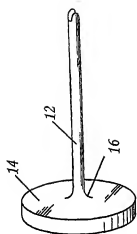


FIG. 1A

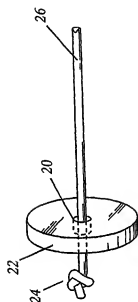


FIG. 1C

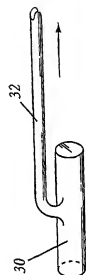


FIG. 1E

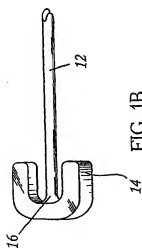


FIG. 1B

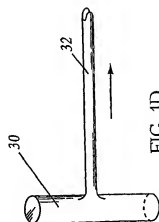


FIG. 1D

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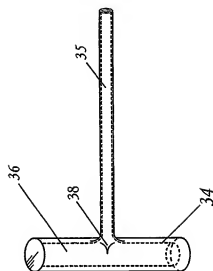


FIG. 1G

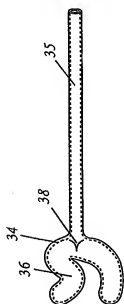


FIG. 1F

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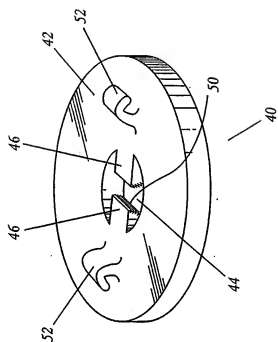


FIG. 2A

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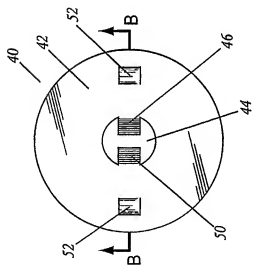


FIG. 2B

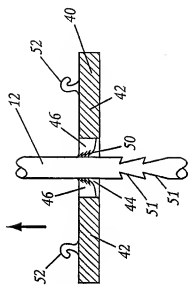


FIG. 2C

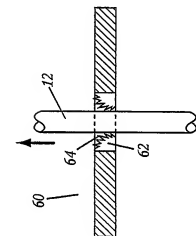


FIG. 2D

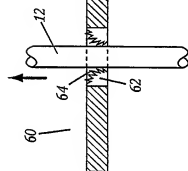


FIG. 2E

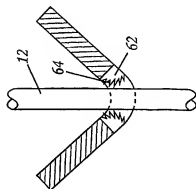


FIG. 2F

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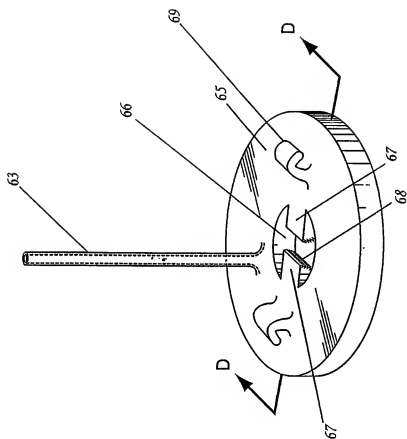


FIG. 2C

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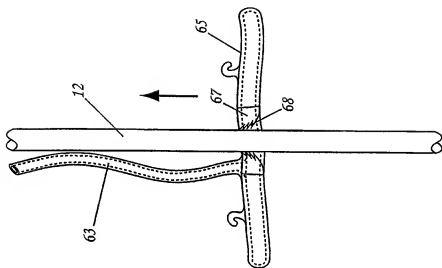


FIG. 2H

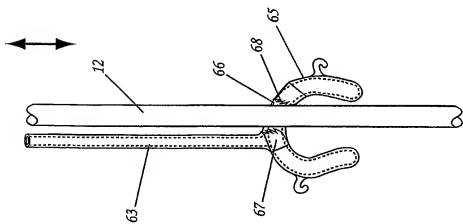


FIG. 2I

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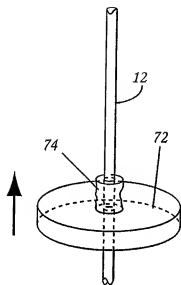


FIG. 3C

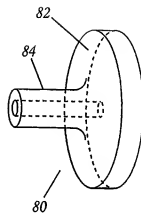


FIG. 3D

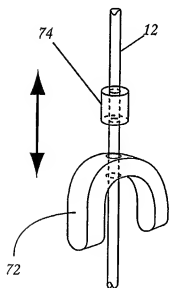


FIG. 3B

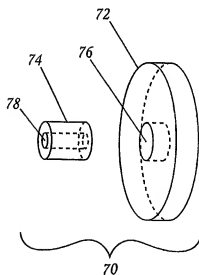


FIG. 3A

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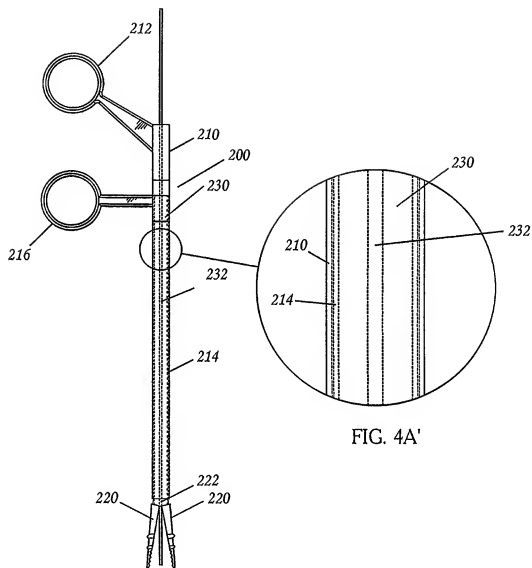


FIG. 4A

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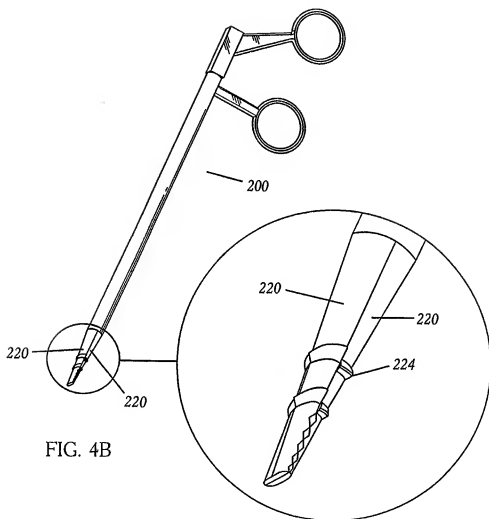
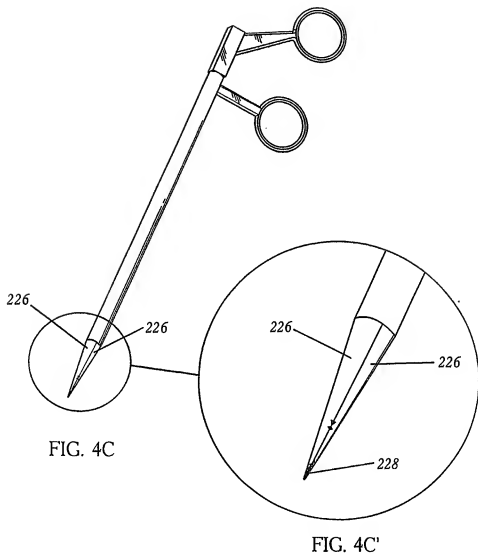


FIG. 4B

FIG. 4B'

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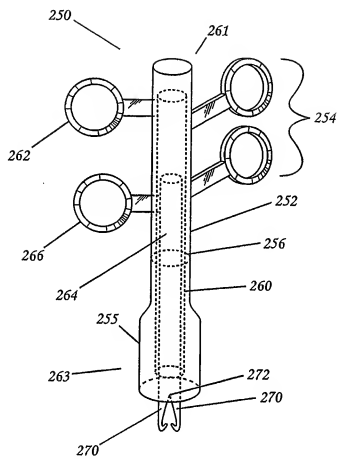


FIG. 5A

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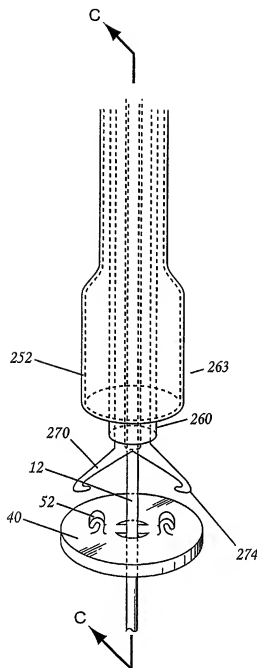


FIG. 5B

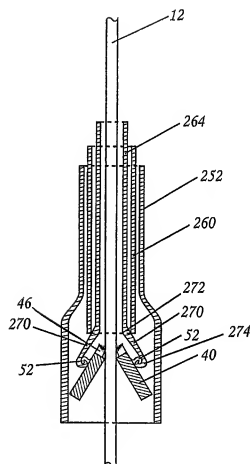


FIG. 5C

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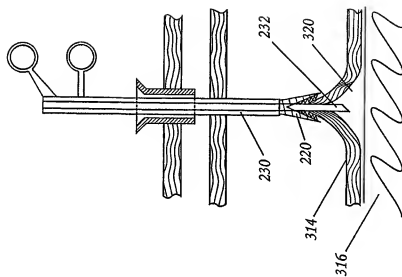


FIG. 6B

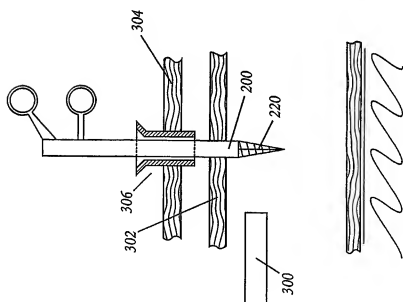


FIG. 6A

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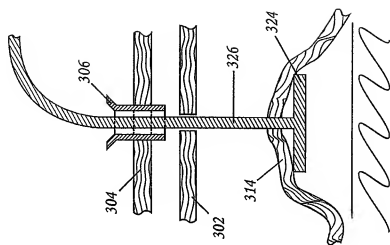


FIG. 6D

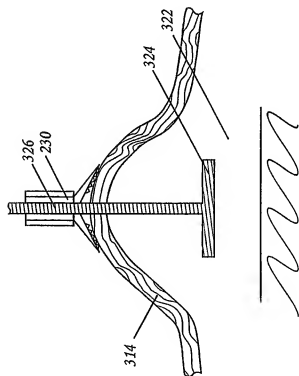


FIG. 6C

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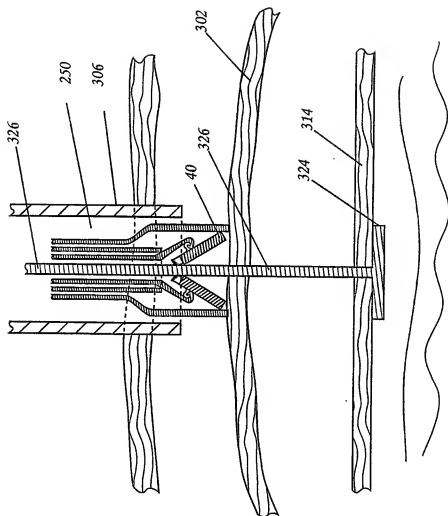


FIG. 7A

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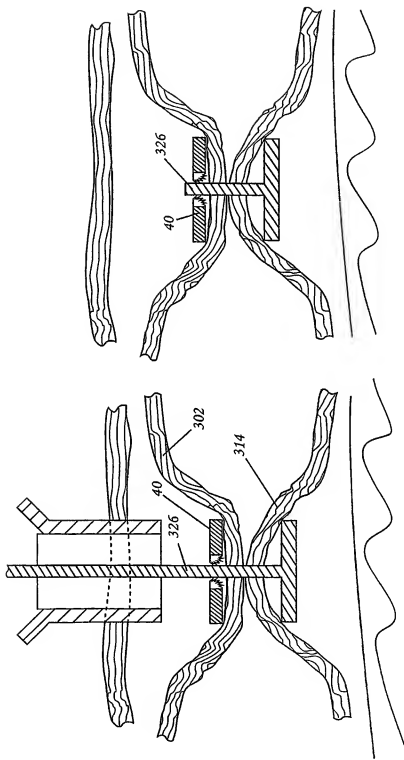


FIG. 7C

FIG. 7B

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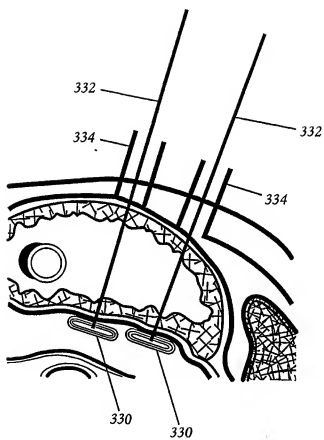


FIG. 8A

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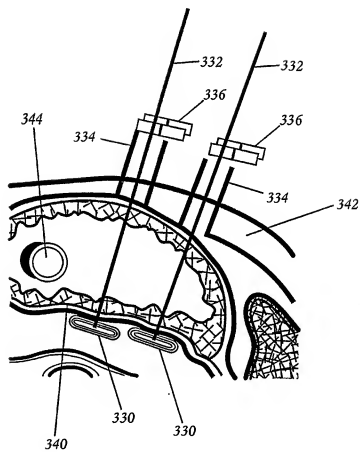


FIG. 8B

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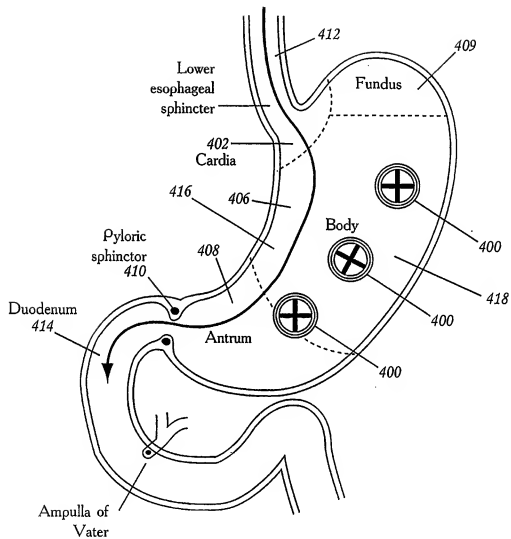


FIG. 9

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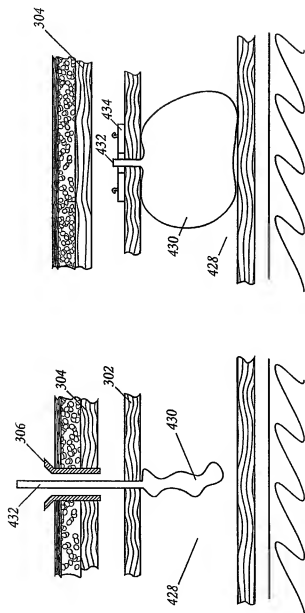


FIG. 10B

FIG. 10A

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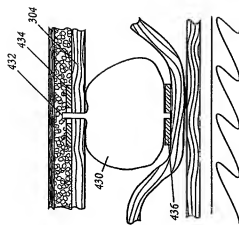


FIG. 11C

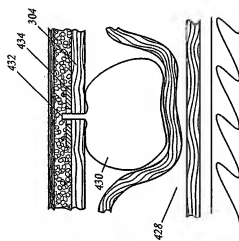


FIG. 11B

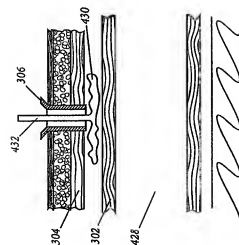


FIG. 11A

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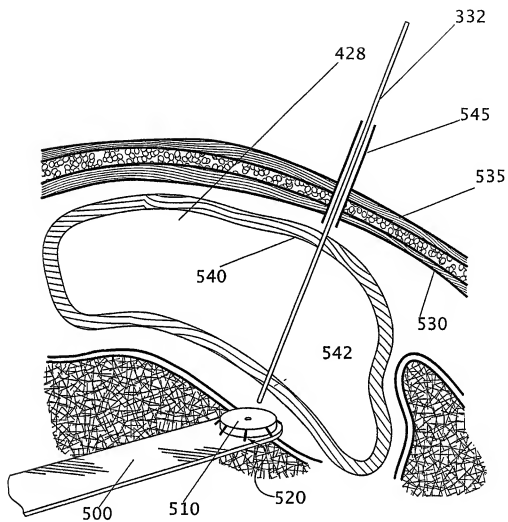


FIG. 12

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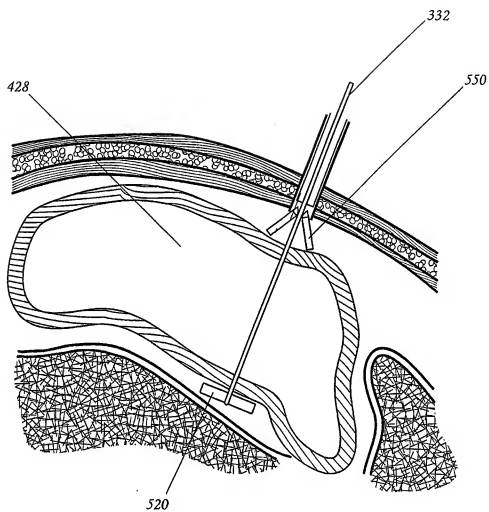


FIG. 13

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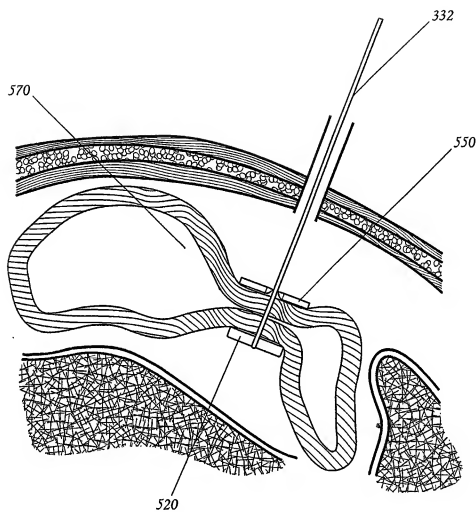


FIG. 14